

GenCore version 4.5
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OM nucleic - nucleic search, using sw model

Run on: July 8, 2002, 12:59:04 ; Search time 5686.68 seconds
(without alignments)
5067.252 Million cell updates/sec

Title: US-09-997-610-3
Perfect score: 1377
Sequence: 1 athgngtncatcngtntc.....gytntatycatcgtatg 1377

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 1797656 seqs, 10463268293 residues

Total number of hits satisfying chosen parameters: 3595312

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%

Listing first 45 summaries

Database :

GenEmbl:*
1: gb_ba:*
2: gb_htg:*
3: gb_in:*
4: gb_lm:*
5: gb_ov:*
6: gb_pat:*
7: gb_ph:*
8: gb_pl:*
9: gb_pr:*
10: gb_ro:*
11: gb_sts:*
12: gb_sy:*
13: gb_un:*
14: gb_vl:*
15: em_ba:*
16: em_fun:*
17: em_hum:*
18: em_in:*
19: em_mu:*
20: em_om:*
21: em_or:*
22: em_ov:*
23: em_pat:*
24: em_ph:*
25: em_pl:*
26: em_ro:*
27: em_sts:*
28: em_un:*
29: em_vl:*
30: em_htg_hum:*
31: em_htg_inv:*
32: em_htg_other:*
33: em_htgo_inv:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Query Match	Length	DB ID	Description
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1	921	66.9	145880	9	HS302D9	282198	Human DNA s
2	497.8	36.2	154090	9	AC025577	AC025577	Homo sapi
3	497.8	36.2	186660	2	AC026107	AC026107	Homo sapi
4	493.6	35.8	124518	9	AL138962	AL138962	Human DNA
5	493	35.8	123070	2	AC008799	AC008799	Homo sapi
6	493	35.8	189768	2	AC044889	AC044889	Homo sapi
7	490.4	35.6	198545	2	AC017063	AC017063	Homo sapi
8	488.8	35.5	161575	2	AC025233	AC025233	Homo sapi
9	488.8	35.5	163639	2	AC087500	AC087500	Homo sapi
10	488.8	35.5	156539	2	AC057727	AC057727	Homo sapi
11	488.6	35.5	161842	2	AL391823	AL391823	Homo sapi
12	488.2	35.5	77405	9	AL353634	AL353634	Homo sapi
13	488.2	35.5	124531	9	HSU212C1	HSU212C1	Human DNA
14	487.6	35.4	40714	9	HSU212C1	HSU212C1	Human DNA
15	487.6	35.4	159712	9	AP004219	AP004219	Human DNA s
16	487	35.4	125295	2	AL672061	AL672061	Homo sapi
17	487	35.4	146743	2	AC093588	AC093588	Homo sapi
18	487	35.4	152544	9	CNS05TEJ	AL359232	Human chr
19	487	35.4	166679	9	AC079899	AC079899	Homo sapi
20	487	35.4	174662	2	AC026036	AC026036	Homo sapi
21	487	35.4	176426	9	AC007370	AC007370	Homo sapi
22	487	35.4	203726	2	AC011882	AC011882	Homo sapi
23	486	35.3	139005	2	AF286112	AF286112	Homo sapi
24	486	35.3	16458	2	AC025647	AC025647	Homo sapi
25	486	35.3	165901	9	AP003474	AP003474	Homo sapi
26	486	35.3	171427	9	AC021590	AC021590	Homo sapi
27	485.8	35.3	99084	2	AC026420	AC026420	Homo sapi
28	485.8	35.3	133790	9	AC010280	AC010280	Homo sapi
29	485.8	35.3	150332	9	AC004921	AC004921	Homo sapi
30	485.8	35.3	174293	9	AC008816	AC008816	Homo sapi
31	485.8	35.3	203043	9	AC017093	AC017093	Homo sapi
32	485.4	35.3	131215	9	AC079614	AC079614	Homo sapi
33	485	35.2	107885	9	AC006389	AC006389	Homo sapi
34	484.6	35.2	173480	9	CNS00MBT	AL079343	Human chr
35	483	35.1	164370	9	AC097460	AC097460	Homo sapi
36	482.6	35.0	112515	2	AL139216	AL139216	Human DNA
37	482.6	35.0	142224	2	AC096535	AC096535	Homo sapi
38	482.6	35.0	178535	2	AL354978	AL354978	Homo sapi
39	482	35.0	168502	9	AC091005	AC091005	Homo sapi
40	481.6	35.0	38235	9	AC004559	AC004559	Homo sapi
41	481.4	35.0	234622	2	AC110089	AC110089	Homo sapi
42	479.8	34.8	187883	9	AC008268	AC008268	Homo sapi
43	479.6	34.8	167891	9	AC013439	AC013439	Homo sapi
44	479.2	34.8	73390	9	AC012038	AC012038	Homo sapi
45	479.2	34.8	94730	9	AP000230	AP000230	Homo sapi

ALIGNMENTS

RESULT 1	HS302D9	145880 bp	DNA	linear	PRI 12-DEC-1999
LOCUS	HS302D9	Human DNA sequence from clone RP1-302D9 on chromosome 22	Contains		
DEFINITION	LOCUS	Human DNA sequence from clone RP1-302D9 on chromosome 22	Contains		
ACCESSION	282198	GI:6572207			
VERSION	282198.2	GI:6572207			
KEYWORDS	HTG.				
SOURCE	human.				
ORGANISM	Homo sapiens				
REFERENCE	1 (bases 1 to 145880)				
AUTHORS	Bridgeman, A.				
TITLE	Direct Submission				
JOURNAL	Submitted (08-DEC-1999) Sanger Centre, Hinxton, Cambridgeshire, CB10 1SA, UK. E-mail enquiries: humquerry@sanger.ac.uk				
COMMENT	On Dec 13, 1999 this sequence version replaced gi:3164067. During sequence assembly data is compared from overlapping clones. Where differences are found these are annotated as variations together with a note of the overlapping clone name. Note that the variation annotation may not be found in the sequence submission				

corresponding to the overlapping clone, as we submit sequences with only a small overlap as described above.

This sequence has been finished according to sequence map criteria as follows. An attempt is made to resolve all sequencing problems, such as compressions and repeats, but not necessarily within known annotated human repeat sequence elements (e.g. Alu). Where the sequence is ambiguous, there is an annotation using the 'unsure' feature key.

The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases:

Em: EMBL; Sw: SWISSPROT; Tr: TREMBL; Wp: WORMPEP; Information on the WORMPEP database can be found at

<http://www.sanger.ac.uk/Projects/C.elegans/wormpep> This sequence was generated from part of bacterial clone contigs of human chromosome 22, constructed by the Sanger Centre Chromosome 22 Mapping Group. Further information can be found at

<http://www.sanger.ac.uk/HGP/Chr22>

RP1-30209 is from the library RPC1-1 constructed at the Roswell Park Cancer Institute by the group of Pieter de Jong. For further details see <http://bacpac.med.buffalo.edu/>

VECTOR: pCYPAC2

This sequence is the entire insert of clone RP1-30209 The true left end of clone CNA-282F2 is at 69682 in this sequence. The true right end of clone CNA-415G2 is at 55167 in this sequence.

FEATURES

Source

1..145880

/organism="Homo sapiens"

/db_xref="taxon:9606"

/chromosome="22"

/clone="RP1-30209"

/clone_1lb="RPC1-1"

188..245

/note="MER3 repeat: matches 144..209 of consensus"

246..571

/note="AluX repeat: matches 1..312 of consensus"

572..759

/note="MER3 repeat: matches 1..144 of consensus"

783..933

/note="MER5A repeat: matches 26..187 of consensus"

1033..1336

/note="AluSp repeat: matches 1..299 of consensus"

1450..1583

/note="MIR repeat: matches 24..160 of consensus"

1687..1752

/note="L2 repeat: matches 2593..2661 of consensus"

2350..2660

/note="AluSc repeat: matches 3..309 of consensus"

2684..2981

/note="AluSq repeat: matches 2..300 of consensus"

3323..3343

/note="MTR1 repeat: matches 116..136 of consensus"

3344..3652

/note="AluY repeat: matches 1..309 of consensus"

3653..3928

/note="MTR1 repeat: matches 136..359 of consensus"

3929..4278

/note="MTR1 repeat: matches 3..364 of consensus"

4279..4485

/note="MTR1 repeat: matches 359..568 of consensus"

5073..5176

/note="52 copies 2 mer ct 78 conserved"

5181..5491

/note="AluB repeat: matches 1..311 of consensus"

6369..6485

/note="L2 repeat: matches 2579..2705 of consensus"

6647..6685

/note="MADEL repeat: matches 1..23 of consensus"

6686..6987

/note="AluX repeat: matches 1..302 of consensus"

6988..7036

/note="MADEL repeat: matches 23..77 of consensus"

7482..7754

/note="AluB repeat: matches 9..290 of consensus"

repeat_region

7775..8060

/note="AluU repeat: matches 1..295 of consensus"

repeat_region

8414..8551

/note="L2 repeat: matches 2553..2706 of consensus"

repeat_region

8914..9030

/note="MIR repeat: matches 147..262 of consensus"

repeat_region

9110..9280

/note="MIR repeat: matches 91..262 of consensus"

repeat_region

9283..9412

/note="MIR repeat: matches 15..144 of consensus"

repeat_region

9521..9679

/note="FAM repeat: matches 3..161 of consensus"

repeat_region

9820..10225

/note="MTR repeat: matches 2..425 of consensus"

misc_feature

10678

/note="match: GSS: Em:856592"

misc_feature

10728

/note="match: GSS: Em:AO701466"

misc_feature

10706

/note="match: GSS: Em:AO225495"

repeat_region

10312..10383

/note="MIR repeat: matches 79..150 of consensus"

misc_feature

10718..11310

/note="match: GSS: Em:B14024"

misc_feature

10784..11201

/note="match: GSS: Em:B43656"

repeat_region

11838..11946

/note="MIR repeat: matches 20..137 of consensus"

repeat_region

12174..12445

/note="L2 repeat: matches 1988..2275 of consensus"

repeat_region

12444..12642

/note="MIR repeat: matches 63..241 of consensus"

misc_feature

13017..13369

/note="match: STS: Em:C49301"

repeat_region

13331..13397

/note="MIR repeat: matches 174..244 of consensus"

repeat_region

13398..13698

/note="AluSp repeat: matches 1..302 of consensus"

repeat_region

13699..13810

/note="MIR repeat: matches 76..174 of consensus"

repeat_region

13806..13919

/note="MIR repeat: matches 77..189 of consensus"

repeat_region

13945..14060

/note="MIR repeat: matches 24..142 of consensus"

repeat_region

14061..14367

/note="AluY repeat: matches 1..301 of consensus"

repeat_region

14368..14452

/note="MIR repeat: matches 141..225 of consensus"

repeat_region

14589..14679

/note="MIR repeat: matches 173..262 of consensus"

misc_feature

14597..15201

/note="match: GSS: Em:AO553482"

misc_feature

14616..15060

/note="match: GSS: Em:AQ370601"

repeat_region

14868..15040

/note="MIR repeat: matches 49..233 of consensus"

repeat_region

15071..15188

/note="L2 repeat: matches 2112..2239 of consensus"

repeat_region

15304..15399

/note="MTR1 repeat: matches 1..99 of consensus"

repeat_region

15490..15662

/note="AluSq repeat: matches 2..114 of consensus"

repeat_region

15669..15727

/note="MTR1B repeat: matches 119..178 of consensus"

repeat_region

15728..16027

/note="AluSc repeat: matches 1..299 of consensus"

repeat_region

16028..16245

/note="MTR1B repeat: matches 178..390 of consensus"

repeat_region

16546..16854

/note="AluY repeat: matches 1..300 of consensus"

repeat_region

18296..18323

/note="MTR1A repeat: matches 2..29 of consensus"

repeat_region

18324..18392

```

/note=MER66-internal repeat: matches 4919. .4993 of consensus"
repeat_region
18393. .18712
/note="AluJb repeat: matches 1. .311 of consensus"
repeat_region
18713. .19133
/note=MER66-internal repeat: matches 4548. .4919 of consensus"
misc_feature
complement(18872. .19230)
/note="match: GSS: Em:A0005063"
misc_feature
19251. .19719
/note="match: GSS: Em:B14179"
repeat_region
19537. .20290
/note=HEVERF21 repeat: matches 4657. .5784 of consensus"
repeat_region
20317. .20382
/note="33 copies 2 mer ta 68 conserved"
repeat_region
20513. .20666
/note="77 copies 2 mer tt 70 conserved"
repeat_region
20682. .21008
/note="Alusg1 repeat: matches 1. .306 of consensus"
repeat_region
21239. .21553
/note="HUMRS-P3 repeat: matches 4410. .4713 of consensus"
repeat_region
21882. .22254
/note="THEIB repeat: matches 1. .364 of consensus"
repeat_region
22302. .22537
/note=MER66-internal repeat: matches 2186. .2417 of consensus"
repeat_region
22538. .22850
/note="Alusp repeat: matches 1. .313 of consensus"
repeat_region
22851. .23801
/note=MER66-internal repeat: matches 1210. .2186 of consensus"
repeat_region
23905. .23989
```

Query Match	66.9%;	Score 921;	DB 9;	Length 145880;
Best Local Similarity	58.2%;	Pred. No. 2e-211;		
Matches 750;	Conservative 300;	Mismatches 239;	Indels 0;	Gaps 0;

OY	89	lbgncncncgngcncncngnyncncncarhtlyacngngarathwtsngaratgcna	148
Db	37254	TGCAATGTCCTGGGTGCCAGGTTTACACAAATATACAGGGAATTAAGTAATACACA	37313
OY	149	atcgcncctcgcncgatalhgarlmwngscntlyacnctgataaryltnwsgnaarytnc	208
Db	37314	AATGCCCTGTCCTGATATAGAAAAGTCAGCCTTACTGTAAGCTCAGTGAACCTTC	37372
OY	209	cnylncntlyacnathahhtlyacngngcnylntlyaycncatmgngayttna	268
Db	37374	CTCTTCCTTTCAAGCCCATCATCTTCAACAGGGCTCTGTCAAAAGCCCAAGGAAATTTA	37433
OY	269	argargcnaatggnglntlycnclymgnglncncngnaaytlaytawsmwntlyag	328
Db	37434	AGAGAGCCATGGAGACTTTCCTTCAAGGGCTCGGGAATTTACTACTCAGCTTTGANG	37493
OY	329	lmgaryltnaayaygygargfnaayahthtgytlnatgmngnaarathytngcnaay	388
Db	37494	TTAGCTGCATCATCTTCAAGGGAATTTTGGCTTAATGAGGAACCAATTTTGGCTTAATA	37553
OY	389	argargaraethwsnarcacarcarwathcargargtnacitggtlytlnylnaargnt	448
Db	37554	AGGAAGAATTTCTTAAGCAGCAACCAATTCATAAGAGGTGACTTGGTCTGTTAAAGCANT	37613
OY	449	lywsntlytlmgngargcngarccayaarwswngaraayltncaycncngayaayttna	508
Db	37614	TCAGTTTCATTAAGGGGAGGAGGCACTTAAGACTTCAGAAAATTTGCCACCCGACAAATGTGA	37673
OY	509	thaaaraaraaaycnclytwsngargngaaatlyaaarytngcngcngarathtlyatht	568
Db	37674	TAAAAAAGAAAACCATTTTCTTGAGGGGAATTCACAGCTGCGCAGAAATTTGCATAT	37733
OY	569	gyaayargaryltnaayltnaayccncargayaaygungnaraayathwtsntgacntgyc	628
Db	37734	GTAAATAGAGAGCTGATATTATTCCTCAAGCAATGGGAAATAATATCTCCTGGACATGCTC	37793

OY	629	armgmwsmwsmcraarwsnatharwsnyhcnctlgmgmccmmgmnaarfbgtcyt	688
Dd	37794	AGAGGCTCTTCACAGAGTCCAAATACACTGGCCGTGAAGGCTTAGGAGAAATGGTTTT	37853
OY	689	gygnaacngmccngmsnynltygtvgtlncarccmmgnaaytngtncnltvgtnc	748
Dd	37854	GTGGAGCAGGCCAGGGTCCCTGTCTGTGTGTCACACCTTAGAGACTTGGTGCCCTGTGTCC	37913
OY	749	cnqlnaaywsngcngtngcnswnsngcngcwnsccnaarcnltvgyarYlncnswng	808
Dd	37914	CAGTTAATTTCAGCTGTGGCTTCAGAGGGTGCAAGGCCCAAGCCTTGCGAGCTTCCAACTG	37973
OY	809	gnltngarcngtngngcnaaraarwsnmgnatlgarYlntvgyarccnccnahtngnt	868
Dd	37974	GGTTTGACCCCTGTGGTGCAAAGACTCAGAAATTTGGGTAACCTCCAAATCAGAT	38033
OY	869	tycaataarathaygnaaycngtltgagtcgcmgmcaarcttvgcngtngngtngnw	928
Dd	38034	TTCAAGATATATATGAAACCCCTGTGATGCCAGGCAAAAGTTTCTCTTAGGGGGGGT	38093
OY	929	snwshltgmgnacnwsngcnmngtngtncarataaygnaaytngntltvggarcncnc	988
Dd	38094	CCTCATGGAGAAACCTCTGCACAGCTAGTACAAABAAGCAAAATTTGGTGGGAGCCCCAC	38153
OY	989	aymgnltncnwsngngncncnswsnmngncngtltmgmgnwsnccnccnswsnm	1048
Dd	38154	ACAGAGTCCCTCCAGTGGGCTCCATCTAGTAGAGCTGTAGAAAGTGCACCATCTCTCCA	38213
OY	1049	gnyltncaraargmgnwsnscngnysnyhncarYlntcngtncngaraarwsnscngya	1108
Dd	38214	GACTGCAGAGGGTAGATTCACACTGCACCTTCACACATGTGCTGTAATAAATCCACAGACA	38273
OY	1109	cncaatgyrcarccngtlnaarngcngngnatlgarYlntcncntayaraarcnngtngtng	1168
Dd	38274	CTCACTGGCAGCGCTTGAAAGCAGCAGGAGTGGATCTGTACCCCTACAAAAACCGTAGTGG	38333
OY	1169	cngraytlnacnaarcnngtngnatlhathaytlncaYltycayayYlntngayltmgnc	1228
Dd	38334	CAGAGCTGACCCAAAGACCGTGGGAATCTACCTTCATATGTCATGACCTGGAGCGTAGAGAC	38393
OY	1229	aygngntnaarmngnaaycayltvngngcnYlmtngtlygarYltygcnaacngntlymna	1288
Dd	38394	ATGGAGTCAAAAGACATCATTTTGGAGCTTTTAAAGATTTGACAGCCCACTGGATTTCCGA	38453
OY	1289	cnlayatvgmccngtncnynlntYltytlyvgncarctYltyccntlygnaacngcngnt	1348
Dd	38454	CTTATATGGGGGCCCTTACCCTTTTGTTTTGGCCATTTTTCATTTTGGAACTGCCGTAT	38513
OY	1349	tyacncartyytlnayYlncayltyatcY 1377	
Dd	38514	TTACCCATATGCTCTGTACCTCCATTTGTATG 38542	

RESULT	2
AC025577	154090 bp DNA linear PRI 25-AUG-2000
LOCUS	
DEFINITION	Homo sapiens 12 BAC RP11-13C3 (Roswell Park Cancer Institute Human BAC Library) complete sequence.
ACCESSION	AC025577
VERSION	AC025577.15
KEYWORDS	GI:9910028
SOURCE	hmg.
ORGANISM	human.
REFERENCE	Homo sapiens
AUTHORS	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo. 1 (bases 1 to 154090) Muzny,D.M., Adams,C., Adio-Oduola,B., All-oman,F.R., Allen,C., Alsbrooks,S.L., Amarantunge,H.C., Are,J.R., Banks,T., Barbata,J., Belton,J., Blmage,K., Blankenburg,K., Bonnin,D., Bouck,J., Boyle,S., Brileva,M., Brown,E., Brown,M., Bryant,N.P., Buhay,C., Burch,P., Burkett,C., Burrell,K.L., Byrd,N.C., Carion,T.F., Carter,M., Cavazos,S.R., Chacko,J., Chavez,D., Chen,G., Chen,R.,

Chen, Z., Chiu, D., Chowdhry, I., Christopoulos, C., Cleveland, C.D., Cox, C., Coyle, M.D., Dathorne, S.R., David, R., Davila, M.L., Davis, C., Davy-Carroll, L., Dederich, D.A., Delaney, K.R., Delgado, O., Dean, A.L., Ding, Y., Dinh, H.H., Douhaite, K.J., Draper, H., Dugan-Rocha, S., Durbin, K.J., Earnhart, C., Edgar, D., Edwards, C.C., Elhaj, C., Emerling, S., Escotto, M., Falls, T., Ferraguto, D., Flagg, N., Ford, J., Foster, P., Frantz, P., Gabisi, A., Gao, J., Garcia, A., Garner, T., Garza, N., Gill, R., Gorrell, J.H., Guayana, W., Gunaratne, P., Hale, S., Hamilton, K., Han, J., Harris, C., Harris, K., Hart, M., Havlak, P., Hawes, A., Hernandez, J., Hernandez, O., Hodgson, A., Hoques, M., Holloway, C., Hollins, B., Homs, F., Howard, S., Huber, J., Hulyk, S., Hume, J., Ioshikhes, I., Jackson, L.E., Jacobson, B., Jia, Y., Johnson, R., Jolivet, S., Joudah, S., Karlsson, E., Kelly, S., Khan, U., King, L., Korvan, J., Kovar, C., Kratovic, J., Kureshi, A., Landry, N., Leal, B., Lee, E., Lewis, L.C., Lewis, L., Li, J., Li, Z., Lichteage, O., Lieu, C., Liu, J., Liu, W., Lounsege, H., Lozano, R.J., Lu, X., Lucier, A., Lucier, R., Luna, R., M.J., Meshwari, M., Mapua, P., Marondel, I., Martin, R., Martindale, A., Martinez, E., Massey, E., Mawhney, E., McLeod, M.P., Meador, M., Mel, G., Merscher, S., Metzger, M., Miller, A., Miner, G., Miner, Z., Mitchell, T., Mohabbat, K., Montgomery, K.T., Morgan, J., Morris, S., Moser, M., Muzny, D., Neal, D., Nelson, D., Newton, M., Newton, N., Nguyen, A., Nguyen, N., Nguyen, N., Nickerson, E., Nwokwkw, S., Ogub, M., Okwuonu, G., Oragunye, N., Oviedo, R., Pace, A., Payton, B., Peery, J., Perez, L., Peters, L., Pickens, R., Primus, E., Pu, L., Quiles, M., Ren, Y., Rives, M., Rojas, A., Rojdoakan, I., Rolfe, M., Ruiz, S., Saverly, G., Scherer, S., Scott, G., Shen, H., Shih, C., Shooshari, N., Sisson, I., Sodergren, E., Sonalike, T., Sparks, A., Stanley, H., Stone, H., Sutton, A., Swatek, A., Taber, P., Telford, B., Thomas, N., Thomas, S., Usmani, K., Vasquez, L., Vera, V., Villalon, D., Vinson, R., Wall, R., Wang, S., Ward-Moore, S., Warren, R., Washington, C., Watlington, S., Williams, G., Williamson, A., Wleczek, R., Woodem, S., Worley, K., Wu, Y., Wu, Y.F., Zhou, J., Zorrilla, S., Kucherlapati, R., Nelson, D. and Gibbs, R.

Direct Submission
Unpublished
2 (bases 1 to 154090)
Worley, K.C.
Direct Submission
Submitted (11-MAR-2000) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 154090)
Worley, K.C.
Direct Submission
Submitted (25-AUG-2000) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
On Aug 25, 2000 this sequence version replaced gi:9664948.
INFORMATION: <http://www.hgsc.bcm.tmc.edu/> or email gc-help@bcm.tmc.edu

CLONE LENGTH: This sequence does not necessarily represent the entire insert of this clone. Overlapping regions of clones are only sequenced and submitted once, so the sequence for the remainder of the insert may be found in the record for the adjacent clones. Overlapping clones are noted at the beginning and end of the Features listing.

ANNOTATION OF FEATURES:
STS are identified using ePCR (Genome Res. 7:541-550) searches of a local database that includes entries from dbSTS, GDB, and local mapping efforts.

Repeats are identified using RepeatMasker (A. Smit and P. Green, unpublished) for Human and Mouse sequences.

Genes and Region of sequence similarity are identified by BLAST (Nuc. Acids Res. 25:3389-3402) similarity (expect < 1e-34) to the EST and cDNA sequences. Genes demonstrate at least two exons flanked by consensus splice sites that maintained sequence continuity across the splice junctions. Sequences that are not identical matches are annotated as similar.

SEQUENCING READ COVERAGE: Sequencing is completed to a minimum standard of double strand coverage with a minimum of 2 clones and 2 reads with no ambiguities or 2 chemistries with a minimum of 2 clones and 3 reads with no ambiguities. If the sequence quality for a region does not meet this standard, it will be indicated in the annotation as Low Coverage.

QUALITY OF INDIVIDUAL BASES: This sequence meets stringent quality standards - estimated error rate less than 1 per 10,000 bases. Reports of lowest quality individual bases and measures of base quality are listed below. Description of the metrics can be found at URL: <http://gc.bcm.tmc.edu:8088/quality.info/genbank.annotation.html>.

QUALSTAT-REPORT-----

----- Summary Statistics -----
Contig length: 154090
Phrap values in estimate: 153545
Average error rate (BCM-Phrap estimate): 0.000261755
Fraction of Phrap values less than 40 : 0.0284086
Number of consensus changing edits: 22
Number of N's in consensus : 0

Position	Original+Context	Edited+Context
9347	aacacagacc(n)tttttttt	aacacagacc(t)tttttttt
10135	attggccct(n)tagaagaaga	attggccct(t)tagaagaaga
10135	ctgtccacg(n)aaagatcca	ctgtccacg(t)aaagatcca
10417	agcaagcag(n)ggctacagaa	agcaatgag(t)ggctacagaa
61959	cttactat(n)tgctatctc	cttactat(t)tgctatctc
61960	ttactatn(t)gtttatctca	ttactatn(g)gtttatctca
61961	ttactatn(t)gtttatctca	ttactatn(g)gtttatctca
61962	taacataa(n)taggttttgg	taacataa(a)taggttttgg
61983	caacacaa(n)taggttttgg	caacacaa(a)taggttttgg
61993	ntaggtttg(n)ttactcttc	ntaggtttg(t)ttactcttc
62205	tcattgacc(n)ctgtcatcc	tcattgacc(a)ctgtcatcc
62956	actgcaacc(n)tgctccacag	actgcaacc(c)tgctccacag
63722	ttcacatata(n)cagtaacgta	ttcacatata(a)cagtaacgta
90216	cctagaanaa(n)gaactttct	cctagaanaa(t)gaactttct
90216	gaaanaagac(n)tttctttt	gaaanaagac(t)tttctttt
90228	ttctttta(n)ttttctaat	ttctttta(t)ttttctaat
90236	tatttttt(n)atcttaagg	tatttttt(t)atcttaagg
90443	accttagatg(n)ctctccacag	accttagatg(a)ctctccacag
91405	cacagcttaa(n)caagtgtaaa	cacagcttaa(a)caagtgtaaa
122290	aaaaaagaa(n)gtaagagaaa	aaaaaagaa(a)gtaagagaaa
137090	acagagaag(n)caaaacatc	acagagaag(a)caaaacatc
141632	tgaagccact(n)gaaaagtaat	tgaagccact(t)gaaaagtaat

----- Distribution of Quality < 40 Bases -----

#	Phrap Value Range
10001	5
9001	10
8001	15
7001	20
6001	25
5001	30
4001	35
3001	40
2001	
1001	
01	

FEATURES

Version: 1.01 gxf.
Location/Qualifiers

Jacobson, B., Jia, Y., Johnson, R., Jolivet, S., Joudah, S.,
Karlsson, E., Kelly, S., Khan, U., King, L., Korvah, J., Kovar, C.,
Kratovic, J., Kuresh, A., Landry, N., Leal, B., Lewis, L. C., Lewis, L.,
Li, J., Li, Z., Lichtarge, O., Lieu, C., Liu, J., Liu, W., Louiseged, H.,
Lozadó, R. J., Lu, X., Lucier, A., Lucier, R., Luna, R., Ma, J.,
Maheshwari, M., Mapua, P., Martin, R., Martindale, A., Martinez, E.,
Massey, E., Mawhinney, E., Mcleod, M. P., Meador, M., Mei, G., Metzger, M.,
Miner, G., Miner, Z., Mitchell, T., Monabhat, K., Morgan, M., Morris, S.,
Moser, M., Neal, D., Newton, J., Newton, N., Nguyen, A., Nguyen, N.,
Nguyen, N., Nickerson, E., Nwokwenkwo, S., Ogun, M., Okunodu, G.,
Oragunye, N., Oviedo, R., Pace, A., Payton, B., Peery, J., Perez, L.,
Peters, L., Pickens, R., Primus, E., Pu, L. L., Quiles, M., Ren, Y.,
Rives, M., Rojas, A., Rojibokan, I., Rolfe, M., Ruiz, S., Saverly, G.,
Scherer, S., Scott, G., Shen, H., Shoohtari, N., Sisson, I.,
Sodergren, E., Sonalke, T., Sparks, A., Stanley, H., Stone, H.,
Sutton, A., Swalek, A., Tabor, P., Tamerisa, K., Tamerisa, K., Tang, H.,
Tansley, J., Taylor, C., Taylor, T., Telford, B., Thomas, N., Thomas, S.,
Umanan, K., Vasquez, L., Vera, V., Villalón, D., Vinsón, R., Wall, R.,
Wang, S., Ward-Moore, S., Warren, R., Washington, C., Watlington, S.,
Williams, G., Williamson, A., Wleczyk, R., Wooden, S., Worley, K.,
Wu, C., Wu, Y., Wu, Y. F., Zhou, J., Zorrilla, S., Nelson, D.,
Weinstock, G. and Gibbs, R.

Direct Submission
Unpublished
2 (bases 1 to 186660)
Worley, K. C.

Direct Submission
Submitted (19-MAR-2000) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA

On May 1, 2001 this sequence version replaced gi:13877175.

* NOTE: Estimated insert size may differ from sequence length
 * (see http://www.hgsc.bcm.tmc.edu/docs/Genbank-draft_data.html).
 * NOTE: This is a "working draft" sequence. It currently
 * consists of 1 contigs. The true order of the pieces
 * is not known and their order in this sequence record is
 * arbitrary. Gaps between the contigs are represented as
 * runs of N, but the exact sizes of the gaps are unknown.
 * This record will be updated with the finished sequence
 * as soon as it is available and the accession number will
 * be preserved.
 * 1 186660: contig of 186660 bp in length.

FEATURES	Location/Qualifiers
source	1. .186660

BASE COUNT	57089 a	37346 c	35962 g	56263 t
ORIGIN				

[illegible]


```

LOCUS       AC008799                123070 bp    DNA    linear    HTG_18-JUL-2000
DEFINITION  Homo sapiens chromosome 5 clone CTD-2061E19, WORKING DRAFT.
ACCESSION   AC008799
VERSION     AC008799.4    GI:9256046
KEYWORDS    HTG: HTGS_PHASE2; HTGS_DRAFT.
SOURCE      human.
ORGANISM    Homo sapiens
REFERENCE   Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS     Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
JOURNAL     DOE Joint Genome Institute.
TITLE       Sequencing of Human Chromosome 5
REFERENCE   2 (bases 1 to 123070)
AUTHORS     DOE Joint Genome Institute.
JOURNAL     DOE Joint Genome Institute.
TITLE       Direct SubMISSION
REFERENCE   Submitted (03-AUG-1999) Production Sequencing Facility, DOE Joint
AUTHORS     Genome Institute, 2800 Mitchell Drive, Walnut Creek, CA 94598, USA
JOURNAL     On Jul 18, 2000 this sequence version replaced gi:7709316.
COMMENT     -----Genome Center
            Center: Joint Genome Institute
            Center code: JGI
            Web site: http://www.jgi.doe.gov
            -----
            Project Information
            Center Project Name: 651088
            Center clone name: CTD-H1_2061E19
            -----
            Summary Statistics
            Consensus quality: 116295 bases at least Q40
            Consensus quality: 121288 bases at least Q30
            Consensus quality: 122086 bases at least Q20
            Estimated insert size: 123000; pulse field gel estimation
            Estimated insert size: 123070; sum-of-contigs estimation
            Quality coverage: 6.04 in Q20 bases; pulse field gel estimation
            Quality coverage: 6.05 in Q20 bases; sum-of-contigs estimation.
            NOTE: This is a 'working draft' sequence. It currently
            * consists of 8 contigs. Gaps between the contigs
            * are represented as runs of N. The order of the pieces
            * is believed to be correct as given, however the sizes
            * of the gaps between them are based on estimates that have
            * provided by the submitter.
            * This sequence will be replaced
            * by the finished sequence as soon as it is available and
            * the accession number will be preserved.
            1
            * 9437 9436: contig of 9436 bp in length
            * 9537 26537: contig of 17001 bp in length
            * 26538 26637: gap of unknown length
            * 26638 45958: contig of 19321 bp in length
            * 45959 46059: gap of unknown length
            * 46059 68657: contig of 22599 bp in length
            * 68657 68757: gap of unknown length
            * 68758 72152: contig of 3395 bp in length
            * 72153 72252: gap of unknown length
            * 72253 81069: contig of 8817 bp in length
            * 81070 81169: gap of unknown length
            * 81170 121547: contig of 40378 bp in length
            * 121548 121647: gap of unknown length
            * 121648 123070: contig of 1423 bp in length.
            Location/Qualifiers
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            /db_xref="taxon:9606"
            /chromosome="5"
            /clone_CTD="2061E19"
            /clone_lib="Caltech human BAC library D"
            37763 a 24223 c 23855 g 36528 t 701 others

```

[illegible]

Db 108927 GGACAGCTGATTACCCATGCTGACCCGATTGTA 108966

RESULT 6 AC044889 189768 bp DNA linear HTG 22-MAY-2000
LOCUS AC044889/c
DEFINITION Homo sapiens chromosome 2 clone RP11-792C1 map 2, WORKING DRAFT
SEQUENCE 36 unordered pieces.
AC044889
AC044889 2 GI:8016676
KEYWORDS HTG; HTGS_PHASE1; HTGS_DRAFT.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Cetartaria; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 189768)
AUTHORS Birren, B., Linton, L., Nusbaum, C. and Lander, E.
TITLE Homo sapiens chromosome 2, clone RP11-792C1
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 189768)
AUTHORS Birren, B., Linton, L., Nusbaum, C., Lander, E., Abraham, H., Allen, N.,
Anderson, S., Baldwin, J., Barna, N., Bastien, V., Bedalov, F.,
Boguslavsky, L., Bouckhaert, B., Brown, A., Burkett, G.,
Campopiano, A., Castle, A., Choquet, Y., Colangelo, M., Collins, S.,
Collymore, A., Cooke, P., Dearlano, K., Dewar, K., Diaz, J. S.,
Dodgson, S., Domingo, M., Doyle, M., Ferreira, P., Fitzhugh, W., Gage, D.,
Gallagher, J., Gardina, S., Glendon, S., Goyette, M., Graham, L.,
Grand-Pierre, N., Grant, G., Hagos, B., Heaford, A., Horton, L.,
Howland, J. C., Iliev, I., Johnson, R., Jones, C., Kahn, L., Karakas, A.,
Klein, J., Lacroque, K., Lamazares, R., Landers, T., Lehoczy, J.,
Lavine, R., Liu, G., Liu, G., Locke, K., Macdonald, P., Margolis, N.,
McCarthy, M., McEwan, P., McGurk, A., McKernan, K., McPheeters, R.,
Meldrum, J., Meneses, L., Mihova, T., Miranda, C., Mianga, V., Morrow, J.,
Murphy, T., Naylor, J., Norman, C. H., O'Connor, T., O'Donnell, P.,
O'Neill, D., Oliver, J. M., Oliver, J., Peterson, K., Pierre, N.,
Pisan, C., Pollara, V., Raymond, C., Riley, R., Rogov, P., Rothman, D.,
Roy, A., Santos, R., Schauer, S., Severy, P., Spencer, B.,
Stange-Thomann, N., Stojanovic, N., Subramanian, A., Talamas, J.,
Traflet, S., Theodore, J., Turrell, A., Travers, M., Triggillo, J.,
Vassiliev, H., Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W. J.,
Young, G., Zainoun, J., Zimmer, A. and Zody, M.

TITLE Direct Submission
JOURNAL Submitted (12-APR-2000) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
COMMENT On May 22, 2000 this sequence version replaced gi:7543856.
All repeats were identified using RepeatMasker:
Smith, A. F. A. & Green, P. (1996-1997)
http://ftp.genome.washington.edu/RM/RepeatMasker.html

Center: Whitehead Institute/ MIT Center for Genome Research

Center code: MIBR

Web site: http://www-seq.wi.mit.edu
Contact: sequence_submissions@genome.wi.mit.edu

Project Information

Center project name: L9594

Center clone name: 792.C.1

Summary Statistics

Sequencing vector: M13; M77815; 100% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.960731
Consensus quality: 168749 bases at least Q40
Consensus quality: 180047 bases at least Q30
Consensus quality: 184014 bases at least Q20
Insert size: 180000; agarose-fp
Insert size: 186268; sum-of-ctnigs
Quality coverage: 3.8 in Q20 bases; agarose-fp
Quality coverage: 3.9 in Q20 bases; sum-of-ctnigs

NOTE: This is a 'working draft' sequence. It currently
* consists of 36 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.

* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
1 1015: contig of 1015 bp in length
1016 1115: gap of 100 bp
1116 2284: contig of 1169 bp in length
2285 2384: gap of 100 bp
2385 3898: contig of 1514 bp in length
3899 3998: gap of 100 bp
3999 5259: contig of 1261 bp in length
5260 5359: gap of 100 bp
5360 6912: contig of 1553 bp in length
6913 7012: gap of 100 bp
7013 8145: contig of 1133 bp in length
8146 8245: gap of 100 bp
8246 9694: contig of 1449 bp in length
9695 9794: gap of 100 bp
9795 11004: contig of 1210 bp in length
11005 11104: gap of 100 bp
11105 12907: contig of 1803 bp in length
12908 13007: gap of 100 bp
13008 14916: contig of 1909 bp in length
14917 15016: gap of 100 bp
15017 17114: contig of 2098 bp in length
17115 17214: gap of 100 bp
17215 18527: contig of 1313 bp in length
18528 18627: gap of 100 bp
18628 20561: contig of 1934 bp in length
20562 20661: gap of 100 bp
20662 22484: contig of 1823 bp in length
22485 22584: gap of 100 bp
22585 25287: contig of 2703 bp in length
25288 25387: gap of 100 bp
25388 27603: contig of 2216 bp in length
27604 27703: gap of 100 bp
27704 30020: contig of 2317 bp in length
30021 30120: gap of 100 bp
30121 33472: contig of 3352 bp in length
33473 33572: gap of 100 bp
33573 38084: contig of 4512 bp in length
38085 38184: gap of 100 bp
38185 41069: contig of 2885 bp in length
41070 41169: gap of 100 bp
41170 45266: contig of 4097 bp in length
45267 45366: gap of 100 bp
45367 51092: contig of 5726 bp in length
51093 51192: gap of 100 bp
51193 55262: contig of 4070 bp in length
55263 55362: gap of 100 bp
55363 60982: contig of 5620 bp in length
60983 61082: gap of 100 bp
61083 67628: contig of 6546 bp in length
67629 67728: gap of 100 bp
67729 72690: contig of 4962 bp in length
72691 72790: gap of 100 bp
72791 79286: contig of 6496 bp in length
79287 79386: gap of 100 bp
79387 83451: contig of 4065 bp in length
83452 83551: gap of 100 bp
83552 92170: contig of 8619 bp in length
92171 92270: gap of 100 bp
92271 96333: contig of 4063 bp in length
96334 96433: gap of 100 bp
96434 103218: contig of 6785 bp in length
103219 103318: gap of 100 bp
103319 112553: contig of 9235 bp in length
112554 112653: gap of 100 bp
112654 123239: contig of 10586 bp in length
123240 123339: gap of 100 bp
123340 137921: contig of 15582 bp in length
137922 138021: gap of 100 bp
138022 157913: contig of 19892 bp in length
157914 158013: gap of 100 bp

OY	750	ngnaaynsngcngtngnswngangr-----g	776
Db	121252	AGCCGCTTAGCATGGCTGAAGAGGGCCCAATGTAGAATAGACGTCAAGCTGTGGCTTCAG	121193
OY	777	ngcmwnuccnaacrcntlbgcarytlctncwmsngngtngarcngtngngcncaaraarws	836
Db	121192	AGGGTGCCTTAAGCCTTGCGCATGTTCCACATNGTGCTTGAAGTCTTTAAGGTACACAGATGC	121133
OY	837	nmgnahtgarqtnltvggarccncnclatmgnltycaraarahcteyggnaaycscntlgtat	896
Db	121132	AAGAATTGAGGTTTTGGGAATCTCCACCCTGGATTTTCAGAAAGATGTATGGMAACGCTTGAT	121073
OY	897	gcocmgncaraazrtlysgngtngtngtngtngmsnsmstngngmaacnwngcmngtngt	956
Db	121072	GCCACGCCAGAAAGTTTGCTGTCAGGGGGCAGGGGATTCATCATAGAACACTCTGCTGAGGGCACT	121013
OY	957	ncaxaiaignaaytfnsgntlggararccnccnaaymgltncnswngngnccnwmws	1016
Db	121012	GCAGAGAGGGAATATGTGGGGTGTGAAGCCACCATATGTGAGTCCCTACTCGGGCACCTTTGAG	120953
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Db	120952	TGGACCTCTGGAAGAGGCGCAATATGACCTCCAGACCCACAGAAATGGTATGTCACCATACAG	120893
OY	1077	nytnaaraytlncnngrraarwnsnacngayaacnarigybarcncngfnaaagcngcngg	1136
Db	120892	CTTGACACCGTATGCTGCTGAAGAAAGCGACAGACACTCAACACCGCCCGTGMAAGCAGCCAG	120833
OY	1137	natgavarvngtncntelyaaaracngtngtngcngaryltnaacnaaracngtngnathla	1196
Db	120832	GAGSGAGSCTGTGCCCCCTACMAAGCCACAGGGGTGAGTGGCCCAAGACCATGGGAACCA	120773
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Db	120772	CCTCTTGATCATGACGATGACCTGGATGTAGAGACTGTGAACAGAGATCATATTTGGAGG	120713
OY	1257	nyltmgntlygaybyrgcnaecnngntlytmgnacntlayatgngncngt-ncenyntlyt	1315
Db	120712	TTTTAAAATTGATATTCCTCTTGCGATTTTCGGACCTTGACCTGTATAAGCCCCTTTGTT	120653
OY	1316	tygngcarctyltycscntlygnaecngcngtlyaacncar	1356
Db	120652	TTTGCCCATTTTCTCCCATTTTGGACAGCGTGTATTATACCCAG	120612
RESULT	9		
ACOSR7500		163674 bp DNA linear HTG 10-JAN-2002	
LOCUS		Homo sapiens chromosome 17 clone RP11-333E1 map 17, *** SEQUENCING	
DEFINITION		IN PROGRESS ***, 11 unordered pieces.	
ACCESSION		AC087500	
VERSION		AC087500.4 GI:18104839	
KEYWORDS		HTG; HTGS_PHASE1; HTGS_FULLTOP; HTGS_ACTIVEFIN.	
SOURCE		human.	
ORGANISM		Homo sapiens	
REFERENCE		Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;	
AUTHORS		Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.	
TITLE		1 (bases 1 to 163674)	
JOURNAL		Biren,B., Linton,L., Nusbaum,C. and Lander,E.	
REFERENCE		Homo sapiens chromosome 17, clone RP11-333E1	
AUTHORS		Unpublished	
		2 (bases 1 to 163674)	
		Biren,B., Linton,L., Nusbaum,C., Lander,E., Allen,N., Anderson,S.,	
		Barna,N., Bastien,V., Boguslavsky,I., Boukhalter,B., Brown,A.,	
		Camarata,J., Campopiano,A., Choepel,Y., Colangelo,M., Collins,S.,	
		Collamore,A., Cooke,P., Dearellano,K., Dewar,K., Diaz,J.S.,	
		Dodge,S., Fato,S., Ferreira,P., Fitzhugh,W., Gage,D., Galagan,J.,	
		Gardyna,S., Glnde,S., Goylete,M., Graham,L., Grand-Pierre,N.,	
		Hagos,B., Heaford,A., Horton,L., Hulme,W., Iliev,I., Johnson,R.,	
		Jones,C., Karatas,A., Larocque,K., Lamazars,R., Landers,T.,	
		Lehocky,J., Levine,R., Liu,G., Maclean,C., MacDonald,P.,	
		Marquis,N., Matthews,C., McCarthy,M., McEvan,P., McKernan,K.,	
		McPheters,R., Meldrum,J., Menius,L., Mihova,T., Mlenga,V.,	

```

Murphy,T., Naylor,J., Nguyen,C., Norbu,C., Norman,C.H., O'Connor,T., O'Donnell,P., O'Neill,D., Oliver,J., Peterson,K., Plunkhardt,P., Pierre,N., Pollara,V., Raymond,C., Retta,R., Riebeck,M., Riley,R., Rise,C., Rogov,P., Roman,J., Rosetti,M., Roy,A., Santos,R., Schauer,S., Schuback,R., Seaman,S., Severly,P., Sougnuez,C., Spencer,B., Strange-Thomann,N., Stojanovic,N., Strauss,N., Traversmanian,A., Talmas,J., Tesfaye,S., Theodore,J., Travers,M., Travis,N., Trigilio,J., Vassiliev,H., Viel,R., Vo.A., Wilson,B., Wu,X., Wyman,D., Ye,W.J., Young,G., Zainoun,J., Zemdek,L., Zimmer,A. and Zody,M.
```

```

Direct Submission
Submitted (05-JAN-2001) Whitehead Institute/MIT Center for Genome Research   330 Charles Street, Cambridge, MA 02141, USA
On Jan 10, 2002 this sequence version replaced gi:13560412.
All repeats were identified using RepeatMasker:
Smtl,A.F.A.&Green,P.(1996-1997)
http://ftp.genome.washington.edu/RM/RepeatMasker.html
```

```
-----Genome Center-----
Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIBR
Web site: http://www-seq.wi.mit.edu
Contact: sequence_submissions@genome.wi.mit.edu
-----Project Information-----
Center project name: L12015
Center clone name: 333_E_1
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* NOTE: This is a 'working draft' sequence. It currently consists of 11 contigs. The true order of the pieces * is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as runs of N, but the exact sizes of the gaps are unknown. * This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.
```

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*      1       3134: contig of 3134 bp in length
*          3135 3234: gap of         100 bp
*          3225 27783: contig of 24549 bp in length
*          27784 27883: gap of         100 bp
*          27884 50218: contig of 22335 bp in length
*          50219 50318: gap of         100 bp
*          50319 54526: contig of 4208 bp in length
*          54527 54626: gap of        100 bp
*          54627 60200: contig of 5574 bp in length
*          60201 60300: gap of         100 bp
*          60301 70374: contig of 10074 bp in length
*          70375 70474: gap of         100 bp
*          70475 79596: contig of 9122 bp in length
*          79597 79696: gap of         100 bp
*          79697 86804: contig of 7108 bp in length
*          86805 86904: gap of         100 bp
*          86905 123464: contig of 36560 bp in length
*          123465 123564: gap of         100 bp
*          123565 160329: contig of 36765 bp in length
*          160330 160429: gap of         100 bp
*          160430 163674: contig of 3245 bp in length.
```

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Location/Qualifiers
    1..163674
     /organism="Homo sapiens"
     /db_xref="taxon:9606"
     /chromosome="17"
     /map="17"
     /clone="RP11-333E1"
     /clone_id="RPCI-11 Human Male BAC"
```

```
BASE COUNT      44782 a  37406 c  35496 g  44882 t  1108 others
```

```
ORIGIN
```

```
Query Match:           35.5% Score 488.8; DB 2; Length 163674;
Best Local Similarity 48.5%; Pred. No. 1.5e+106;
```

```
Matches 485; Conservative 202; Mismatches 280; Indels 34; Gaps 2;
```

```
OY 390 rgargaratnnaarcacrcarwcnathcagatglnactgggtunynynaargcnt 449
```

```
[:::|||||:] [|||] : || | ||| || :: || | || | :
```


[illegible]

Db	9477	AGGAGGCGCTGACCTCGCAAGGCCACAGGGGACAGCTGCCCAAGATCATGGAAACCAAT	9536
Oy	1198	yltynltncaeytbycaaygayyltngayyltmngncaaynglnaarmngaycayltygngcn	1257
Db	9537	CTCTGTCATCGACGACCTGGATATGAGATGGAGTCAAGGAATAATTTGGAGCT	9596
Oy	1258	yltmngntlygagyltygcacnngntlymgnacltayaatggngcngtncnynlntaylt	1316
Db	9597	TTAAATATTGACGTGCCACAGGATCTCAGACTTCACAGGGGGCTGTAGCCTCTTTGT	9656
Oy	1317	ygnaarltlytlycngntlygnaacngntlytlyacnartlytlytnaeyltncaeytlat	1376
Db	9657	TGGCCAAATCTCCATTGGAAATGCGTGTATTATCAAAATGCCCTGACCCCAATTGTAT	9716

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RESULT 13
HSJ633H17/c
LOCUS      124531 bp      DNA      linear      PRI 07-FEB-2000
DEFINITION Human DNA sequence from clone 633H17 on chromosome 1p31.2-32.2.
             Contains a pseudogene similar to part of MTCO1 (cytochrome C
             oxidase 1), MTCO2 (cytochrome c oxidase ii), MTATP8 (ATP synthase
             8) and GORT2 (glutamic-oxalacetic transaminase 2, mitochondrial
             (aspartate aminotransferase 2) pseudogenes, ESTs and GSSs, complete
             sequence.
ACCESSION  AL049710      GI:5650682
VERSION     AL049710.18
KEYWORDS    HTG; aspartate aminotransferase; ATP synthase; Cytochrome C
             oxidase; glutamic-oxalacetic transaminase; GORT2; MTATP8; MTCO1;
             MTCO2.
SOURCE      human.
ORGANISM    Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
REFERENCE   1 (bases 1 to 124531)
AUTHORS     Moore,M.
TITLE       Direct Submission
JOURNAL     Submitted (21-SEP-1999) Sanger Centre, Hinxton, Cambridgeshire,
            CB10 1SA, UK. E-mail enquiries: humquerry@sanger.ac.uk
COMMENT     On Jul 29, 1999 this sequence version replaced gi:5566565.
            During sequence assembly data is compared from overlapping clones.
            Where differences are found these are annotated as variations
            together with a note of the overlapping clone name. Note that the
            variation annotation may not be found in the sequence submission
            corresponding to the overlapping clone, as we submit sequences with
            only a small overlap as described above.
            This sequence is the entire insert of clone 633H17. This sequence
            has been finished according to sequence map criteria as follows. An
            attempt is made to resolve all sequencing problems, such as
            compressions and repeats, but not necessarily within known
            annotated human repeat sequence elements (e.g. Alu). Where the
            sequence is ambiguous, there is an annotation using the 'unsure'
            feature key.
            This sequence was generated from part of bacterial clone contigs of
            human chromosome 1, constructed by the Sanger Centre Chromosome 1
            Mapping Group. Further information can be found at
            http://www.sanger.ac.uk/HGP/Chr1
            633H17 is from the library RPC1-4 constructed at the Roswell Park
            Cancer Institute by the group of Pater de Jong. For further
            details see http://bacpac.med.buffalo.edu/VECTOR:pcrPAC2 The
            following abbreviations are used to associate primary accession
            numbers given in the feature table with their source databases:
            Em1, EMBL; Sw1, SWISSPROT; Tr1, TREMBL; Wp1, WORMPEP; information
            on the WORMPEP database can be found at
            http://www.sanger.ac.uk/Projects/C_elegans/wormpep.

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                /db_xref="taxon:9606"
                /chromosome="1"
                /map="p31.2-32.2"
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2026. .2590
repeat_region /note="HERVL repeat: matches 4676. .5241 of consensus"
2879. .9441
repeat_region /note="L1P2 repeat: matches 1. .6146 of consensus"
10000. .10124
repeat_region /note="MIR repeat: matches 34. .154 of consensus"
10623. .10689
repeat_region /note="MIR repeat: matches 36. .102 of consensus"
10690. .11014
repeat_region /note="L2 repeat: matches 2389. .2710 of consensus"
11136. .11262
repeat_region /note="L2 repeat: matches 2349. .2473 of consensus"
13420. .13522
repeat_region /note="MIR repeat: matches 114. .219 of consensus"
13602. .13725
repeat_region /note="MIR repeat: matches 130. .262 of consensus"
13739. .13987
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14368. .14486
repeat_region /note="L2 repeat: matches 2585. .2705 of consensus"
14656. .14764
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14966. .15150
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18377. .18492
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18525. .18954
misc_feature /note="match: GSS: Em:AQ210513"
19026. .19126
repeat_region /note="MIR repeat: matches 10. .114 of consensus"
19866. .20007
repeat_region /note="MIR repeat: matches 28. .169 of consensus"
20273. .20334
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20403. .20505
repeat_region /note="MER81 repeat: matches 1. .110 of consensus"
20508. .20690
repeat_region /note="L1R16C repeat: matches 190. .376 of consensus"
20692. .20725
repeat_region /note="L2 repeat: matches 2660. .2693 of consensus"
20828. .20863
repeat_region /note="18 copies 2 mer ac 94% conserved"
21172. .21304
repeat_region /note="MERSA repeat: matches 40. .167 of consensus"
21201. .21331
repeat_region /note="MERSA repeat: matches 56. .188 of consensus"
21466. .21528
repeat_region /note="MERS4A repeat: matches 214. .285 of consensus"
21952. .22077

repeat_region /note="L2 repeat: matches 2578. .2707 of consensus"
22083. .22259
repeat_region /note="FRAM repeat: matches -2. .175 of consensus"
22280. .22393
repeat_region /note="L2 repeat: matches 2541. .2645 of consensus"
22394. .22748
repeat_region /note="M1R1B repeat: matches 1. .390 of consensus"
22749. .22846
repeat_region /note="L2 repeat: matches 2645. .2745 of consensus"
22864. .23010
repeat_region /note="L2 repeat: matches 2329. .2429 of consensus"
23011. .23310
repeat_region /note="Alu1b repeat: matches 1. .299 of consensus"
23311. .23391
repeat_region /note="L2 repeat: matches 2429. .2521 of consensus"
23433. .23730
repeat_region /note="M1TD repeat: matches 66. .394 of consensus"
23731. .25397
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25477. .25556
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25557. .25651
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26898. .27373
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27484. .27868
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27912. .28051
repeat_region /note="L1P13 repeat: matches 6011. .6152 of consensus"
29900. .30121
repeat_region /note="HERVL repeat: matches 3020. .3235 of consensus"
30122. .30487
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33947. .34302
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34335. .34703
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34704. .34830
repeat_region /note="MIR repeat: matches 20. .129 of consensus"
34831. .35552
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Query Match

35.5%, Score 488.2; DB 9; Length 124531;


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	Query Match	35.4%	Score 487.6;	DB 9;	Length 40714;
	Best Local Similarity	47.0%;	Pred. No. 1.8e-106;		
	Matches 507,	Conservative 225;	Mismatches 311;	Indels 36;	Gaps 4;
OY	334 yncycaytgyaaagttaaayaathhggtynatgmnacarcathlytngcmaayaargar	393			
	: : : : : : : : : : : : : : : : : : : : : :				
Dd	24003 CCGCCCTGAGAGATCTGTGGAAATTTCACCTTGAGAGAGATGATTAGGGTATCAGGTGCA	24062			
	: : : : : : : : : : : : : : : : : : : : : :				
OY	394 garalhwnaacrcarcarsnatlhargarynactggtylrythytaarycnlttysn	453			
	: : : : : : : : : : : : : : : : : : : : : :				
Dd	24063 GAAATTTCTAACACCACAAGCATTCAGAGGGGACTTGTCCTTTAAAGCACTCCAT	24122			
	: : : : : : : : : : : : : : : : : : : : : :				

[illegible]

SOURCE Homo sapiens pre-pro-B cell clone:FLB 14 - 14 DNA,
clone_lib:keio library clone:KB1222D11.

ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.

REFERENCE 1 (bases 1 to 159712)
Shimizu N. and Asakawa, S.

AUTHORS Shimizu N.

TITLE Direct Submission

JOURNAL Submitted (28-SEP-2001) Nobuyoshi Shimizu, Keio University, School
of Medicine, Molecular Biology; 35 Shinanomachi, Shinjuku-ku, Tokyo
160-8582, Japan (E-mail:nshimizu@med.keio.ac.jp,
Tel:81-3-3351-2370, Fax:81-3-3351-2370)

COMMENT On Jan 14, 2002 this sequence version replaced gi:15824050.

FEATURES
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Location/Qualifiers
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/map="8q23"
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/cell_line="FLB 14 - 14"
/cell_type="pre-pro-B cell"
/clone_lib="Keio BAC library"
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repeat_region
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9188..9226
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20567..20636
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/rpt_family="MER94"
repeat_region
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Query Match	35.48:	Score 487.6:	DB: 9:	Length 159712:
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Db 138232	AGAGAAAGTCTCTTAAGCAGCAAGACATTCACAGAGATTACTGGGAGTGTCATTAAAAGCACT	138173		
Qy 450	ywnsttvtvltmgngaragcngaracvayaaatwsmngararaytlnaycncngayaaqytnat	509		
Db 138172	CAGTTTATTAAAGGAGAGAGACATAAAACTTTGGAATAATTCGACGCTGACTATGACAT	138113		
Qy 510	haaraaraaraaaycncntlywsngaragynaartlyaaeryltnngcngarathctyathlytg	569		
Db 138112	AGAAAAGAAAAAACCATTTTCGCGGAGAAATTCACAGCGGCTGCAGAAATTTGCGACAAAG	138053		
Qy 570	yaayaqargarylnaayqylnaayccmcarqargayaaayngngararayathwntgaactlytca	629		
Db 138052	TAGCAAGAGCCAAATTTTAATCCCAAGACCATGGGAAAATGTCCTCAGGCCATTGCA	137993		
Qy 630	rmngwsmnsncarcarwsnathaarwsmlytngcnltgmgnc--cmmgmngmaarttqglt	686		
Db 137992	GAGACCTTATAGCCAGCCCTCCATCTATTAAGCTCGAGAGCCTACGAGAAAATGGTGT	137933		

QY	687	yttygnaengnccngnswsnynttytgytlnearccmmgayaftnglncnhtygt	746
Db	137932	TCATGGGCTGGCCAGGGCTCCCTGGGGCTGTGTGCACGCTTAGGAACTGGTGTGCACCTGTGT	137873
QY	747	ncngtnaaysnscngtngcnwngangr-----ncnwsnccnaa	788
Db	137872	CCCAACCCACTCAGCCGGCTGAAAGGGGCGCACTGTACACCTTGGCTGTGGCTTCACA	137813
QY	789	rcnhtygcarytncnwsnsgngtngearccngtngngcngnaaraazsmnaglnhgyrt	848
Db	137812	GGCTTGGCAGCTTCACACTGGTGTGTAAGCCCTGTGGGTGGCACAAGACGACAAAGTAATGAGCT	137533
QY	849	nltygarcncncnaltmgnttycacaarathlaygnaaycnclygatlgccmmncaraa	908
Db	137752	TTGGGAACCTCTGCCTAGATTTCAGAAGATGTATGAAATATCCCTGATGGCCAGGCCAAAA	137693
QY	909	rttycngtngtngtngnswsnwntltvgmgnacnwsngcmngtnglncaraargnaa	968
Db	137692	GTTTCTCTCAGAGGGGGGGCCCTCATGGAGAACTCTGTCTTAGGGCACTGAGAAAGAAAA	137633
QY	969	ytngngtngygarccnccncaymngltnccnwsngngnccnwsnsmngcngtnglmg	1028
Db	137632	TGTGGGGTTCGATTTCACACACAGAGTCCCTACTGCGGGCACGTGCTAGTGGAGCTGTGAG	137573
QY	1029	nmgnwsnccnccnswsnwmngnylncaaraargmngmgnasnagnayaasnylncaarcayt	1088
Db	137572	AAGAGGCGACACATCCTTCAGAACCCCAAGAAATGTATCCACTGCACAGCTTGACACCATGT	137513
QY	1089	ncnngaraarasnagnayaencarttycarccngtlnaargcngcngnagatgtarwngt	1148
Db	137512	GCCTTGAAAAAATCGAGACATTCAATGCGACACCCATGAAGAAGCACGCCGAGAGAGGCTAT	137453
QY	1149	ncnhtayaarznngtngtngcngarythncaaraacngtnglnaathaytlnyngtyt	1208
Db	137452	ACCCTGCAAAAGCACAGGGGAGAGACTCCCAAAACATATGAAACCTTCTTGTGCACTCA	137393
QY	1209	ycaaygaytlnagytlnmncacaygngtlnaaraargnayaaytlytgngcngnylnmgntlyga	1268
Db	137392	GGGTACCTCGATGTGAAACCTGTATATCAAGAAGATCATTTTGGACCTTTAAATTTGG	137333
QY	1269	ytyygcnaengntlytmnacinayatlvgngcngt--ncnynlttytlyvgncarttyt	1327
Db	137332	CTGGCCCACTGTTTGTGGACTTCATCCATGGGCCCTGTAAACCTCTTGTGTGGCAATATTT	137272
QY	1328	tycncntlygnaengcngtlnthyacncarttyytnataytlncaityat	1376
Db	137272	TCCCAATTTTGGAAACATAGCTGTTTCCCAATACCTGTATCTCCCAATTTGAT	137224

Search completed: July 8, 2002, 18:08:58
Job time: 18594 sec

.....

GenCore version 4.5
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OM nucleic - nucleic search, using sw model

Run on: July 8, 2002, 12:53:47 ; Search time 401.64 Seconds

(without alignments)
4908.627 Million cell updates/sec

Title: US-09-997-610-3

Perfect score: 1377

Sequence: 1 atggtgncatcncgntgnt.....gyttntaytccaytgyatg 1377

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 1736436 seqs, 858457221 residues

Total number of hits satisfying chosen parameters: 3472872

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :
1: N_Geneseq_032802:*
2: /SID5/gcgcdata/geneseq/geneseqn-emb1/NA1980.DAT:*
3: /SID5/gcgcdata/geneseq/geneseqn-emb1/NA1981.DAT:*
4: /SID5/gcgcdata/geneseq/geneseqn-emb1/NA1983.DAT:*
5: /SID5/gcgcdata/geneseq/geneseqn-emb1/NA1984.DAT:*
6: /SID5/gcgcdata/geneseq/geneseqn-emb1/NA1985.DAT:*
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23: /SID5/gcgcdata/geneseq/geneseqn-emb1/NA2001B.DAT:*
24: /SID5/gcgcdata/geneseq/geneseqn-emb1/NA2002.DAT:*

pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	471.4	34.2	2590	22	AAH14327
2	452	32.8	3093	22	AA503687
3	452	32.8	9236	22	AA503689
4	452	32.8	9236	22	AA503690
5	452	32.8	9241	22	AA503698
6	429.8	31.2	3203	22	AA005134
7	427.6	31.1	6063	22	AA003635
8	427.2	31.0	22509	22	AA544505
9	427.2	31.0	31529	22	AA544506

10	426.4	31.0	6063	22	AA03634	Human reproductive
11	421.4	30.6	1736	22	AAH13678	Human CDNA sequenc
12	421	30.6	465237	22	AAH13678	Human oestrogen re
13	409	29.7	1485	23	AA591984	DNA encoding novel
14	408.8	29.7	1278	22	AA005178	Human secreted pro
15	400.4	29.1	1580	23	AA587262	DNA encoding novel
16	395.2	28.7	7726	22	ABA21079	Human nervous syst
17	393.6	28.6	32127	22	AAI92955	Human excretory re
18	393.6	28.6	32127	22	AAI63605	Human kidney relat
19	393.6	28.6	72215	22	AAK68632	Human immune/haema
20	388.2	28.2	1405	22	AAK70265	Human immune/haema
21	386.6	28.1	1405	22	AAK70266	Human immune/haema
22	386.4	28.1	8387	22	ABA14892	Human nervous syst
23	386.4	28.1	8387	22	ABA84548	Human immune/haema
24	384.4	27.9	1621	23	AA567924	DNA encoding novel
25	378.8	27.5	3049	22	AAH16637	Human CDNA sequenc
26	373.8	27.1	81369	21	AA57997	Human T gene DNA.
27	371.6	27.0	1306	23	AA564678	DNA encoding novel
28	361.6	26.3	22756	22	AA540321	DNA encoding human
29	361.6	26.3	22756	22	AA03921	Human reproductive
30	355.4	25.8	1946	22	ABA44227	Human breast cell
31	355.4	25.8	1946	22	ABA54677	Human foetal liver
32	355.4	25.8	1946	22	ABA24459	Probe #2925 for ge
33	355.4	25.8	1946	22	AAK02964	Human brain expres
34	355.4	25.8	1946	22	AAK84410	Human bone marrow
35	355.4	25.8	1946	22	AAI12975	Probe #2908 for ge
36	355.4	25.8	1946	22	AAI34334	Human T cell
37	355.4	25.8	1946	22	AAI02893	Human foetal liver
38	347	25.2	26410	22	AAK70623	Human immune/haema
39	346.8	25.2	32986	22	AAK69758	Human immune/haema
40	346.8	25.2	32986	22	AAK69758	Human immune/haema
41	346.4	25.2	32986	22	AAK69758	Human immune/haema
42	341.4	24.8	50000	24	AA026400	DNA encoding novel
43	341.4	24.8	50000	24	AA026437	Human GRM3 gene fr
44	337.4	24.5	57728	22	AA087588	Human p11 chromos
45	336.6	24.4	2197	22	AAH18389	Human CDNA sequenc

ALIGNMENTS

RESULT 1	
ID	AAH14327 standard; CDNA; 2590 Bp.
XX	AAH14327:
AC	AAH14327:
XX	26-JUN-2001 (first entry)
DT	26-JUN-2001 (first entry)
XX	Human CDNA sequence SFG ID NO:11697.
DE	Human CDNA sequence SFG ID NO:11697.
XX	Human; primer; detection; diagnosis; antisense therapy; gene therapy; ss.
KW	Human; primer; detection; diagnosis; antisense therapy; gene therapy; ss.
XX	Homo sapiens.
OS	Homo sapiens.
XX	EP1074617-A2.
PN	EP1074617-A2.
XX	07-FEB-2001.
PD	07-FEB-2001.
XX	28-JUL-2000; 2000EP-0116126.
PF	28-JUL-2000; 2000EP-0116126.
XX	29-JUL-1999; 99JP-0248036.
PR	29-JUL-1999; 99JP-0248036.
XX	11-JAN-2000; 2000JP-0118776.
PR	11-JAN-2000; 2000JP-0118776.
XX	02-MAY-2000; 2000JP-0183767.
PR	02-MAY-2000; 2000JP-0183767.
XX	09-JUN-2000; 2000JP-0241899.
XX	(HELI-) HELIX RES INSR.
PA	(HELI-) HELIX RES INSR.
XX	Ota T, Isogai T, Nishikawa T, Hayashi K, Saito K, Yamamoto J;
PI	Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;
XX	WPI; 2001-318749/34.

Pt	Primer sets for synthesizing polynucleotides, particularly the 5602
Ft	full-length cDNAs defined in the specification, and for the detection
Pt	and/or diagnosis of the abnormality of the proteins encoded by the
Pt	full-length cDNAs -
Pt	xx
Ps	Claim 8; SEQ ID 11697; 2537bp + CD ROM; English.
Cc	The present invention describes primer sets for synthesising 5602
Cc	full-length cDNAs defined in the specification. Where a primer set
Cc	comprises: (a) an oligo-dT primer and an oligonucleotide complementary
Cc	to the complementary strand of a polynucleotide which comprises one of
Cc	the 5602 nucleotide sequences defined in the specification, where the
Cc	oligonucleotide comprises at least 15 nucleotides; or (b) a combination
Cc	of an oligonucleotide comprising a sequence complementary to the
Cc	complementary strand of a polynucleotide which comprises a 5'-end
Cc	sequence and an oligonucleotide comprising a sequence complementary to a
Cc	polynucleotide which comprises a 3'-end sequence, where the
Cc	oligonucleotide comprises at least 15 nucleotides and the combination of
Cc	the 5'-end sequence/3'-end sequence is selected from those defined in
Cc	the specification. The primer sets can be used in antisense therapy and
Cc	in gene therapy. The primers are useful for synthesising polynucleotides,
Cc	particularly full-length cDNAs. The primers are also useful for the
Cc	detection and/or diagnosis of the abnormality of the proteins encoded by
Cc	the full-length cDNAs. The primers allow obtaining of the full-length
Cc	cDNAs easily without any specialised methods. AAH03166 to AAH13628 and
Cc	AAH13632 to AAH18742 represent human cDNA sequences; AAB92446 to
Cc	AAB95893 represent human amino acid sequences; and AAH13629 to AAH13632
Cc	represent oligonucleotides, all of which are used in the exemplification
Cc	of the present invention.
Cx	xx
SQ	Sequence 2590 BP; 722 A; 526 C; 670 G; 672 T; 0 other:
Query Match	34.2%; Score 471.4; DB 22; Length 2590;
Best Local Similarity	47.0%; Pred. No. 1.1e-108;
Matches	481; Conservative 213; Mismatches 293; Indels 37; Gaps
OY	360 rgaargathtwsnaarcacarnasathcaargatnactntgygtintylnaarcntt 449
Db	: : :: : :: : : :: :
OY	608 agaatattcttaagccagaacaagtccaagaagttagtgttgcatctgttaagaacct 667
OY	450 ysaanttyatlmngnrgncrcarcalyaatwswnsngnaayrincycocngaayaatnat 509
Db	:: :: : :: :: :: :: : :: :
OY	668 cgatttcataaaagggaacaaagcataaaaattcagaanaatttgttagcctgcatacgtaat 727
OY	510 haaraaraaraycnctlytsngartggnaartltyaarlyungonggnarratltyatbtg 569
Db	: :: :: : :: :: :: :: :: :
OY	728 agaanaaaaaaacccagtttctcgtgggagaaattcaaaccagcgtcgaagaatactgcataag 787
OY	570 yaayaargaralytnaagythaacyccnacargayaagyngnaryaalathwsnttgacntlgyca 629
Db	: : : : :: :: : :: :: : : :
OY	788 catcaagaagaccttaatgtlaattcccacagcacatlgvggaatatgtctccagcattgca 847
OY	630 tmgwnsnscnarcarwsnathtaarsnylungentgymgncmngm---naartggtt 686
Db	:: : :: : : :: : : : : : :
OY	848 gagaccttcacagcagccccctctctgltcaacagccccagagatccagagagaaataatggtt 907
OY	687 ycygysnagnngncongngwnsyntlytygytlncarcmngnangrynlngntncttygt 746
Db	: :: : :: :: : :: :: :
OY	908 ttatygccacaggaaccaagggtccctctgctgtgltgcaagctaataagacttgggtccctgtgt 967
OY	747 nccngtlnaawys-----ngcgttgcmwsnga 773
Db	: : : : : :: : :
OY	968 ccagcactgctccagccatcagctgataaaggggccaatgtagagctatctgtctgcttcaga 1027
OY	774 rgngcnwnscnnaarcncttggarlytncnmwsgnngtlnagarcnngtngngncaaraa 833
Db	: :: : :: :: : :: :: :
OY	1028 ggggtggaagccccaagccttgagcagcttcacatgltgttgagcctgtgagfcaacagaa 1087
OY	834 twsmgmnahtargtntgggartccmcnalmngtlycaraarthtaygnaaycentg 893
Db	: : :: :: :: :: :: : :: :: :
OY	1088 gccaggaatttgaggtttggaaacctccacactgatatttcagaagaatglatgtaaagctgct 1147

Oy	894	gacgccmngncararatttgcgctgngtngngtngmwnsnbtgmgbnacnwnsgcmngt	953
Db	1148	gatctccgagcaaaagtcttgcacagtggcaaggcccttataggaaaccttcgctaaaggc	1207
Oy	954	ngtncaraagnaaygttngtgnltggarcnccncaymngtncnccmwnsgngnccmws	1013
Db	1208	aatgcagaagggaattctgtgggcccggagctcccaagctagagtcctaatggygcatttc	1267
Oy	1014	nwsnmngcngltmngmgnwnsnccnccmwnsmngnyltncaraarngmngwnsnacnga	1073
Db	1268	tagtggagctgtggagaagaggccacgcgtcccttcagaccacccagcatgtgtgatacctga	1327
Oy	1074	ywsnlytncaarccygtncncngaraatwsmnccngayacncaarttgcacrcctngtaargcnc	1133
Db	1328	caacttgcacacatgycgcgcggaaagccgcgacacactcaatgcagtcctgtgaaagcagc	1387
Oy	1134	ngnatctgarwngtncnccntaaraacngtngtngcnarytnacnaaracngtngnat	1193
Db	1388	caggagggagagctgttacccttgaaaggcccaaggcagagacgcgccaaaggccatgggtcc	1447
Oy	1194	htaytlytncaaycgycaaygayttnagytltmngnacayggngtlnaarmngaycaytlygg	1253
Db	1448	ccacctttgcacacagctgcacactgaattgagacctaagtcacaaagagacatcatttgg	1507
Oy	1254	ngcyltmngnttgyaytgyccnaccngnttmgnaentatayatzggncng - tncnnynt	1312
Db	1508	agctctaaaatctgtatctgcgcgcggatcttgagacttgcaggggtccctgcagcccttc	1567
Oy	1313	gytlygncarttlytccnttlyggnacngcngtnttcaacncaarttgytntaytlncayt	1372
Db	1568	gttttgcacatgctcccatctgaatgtgcgtatattaccataacctgtaccctatcc	1627
Oy	1373	gyat	1376
Db	1628	great	1631
RESULT 2			
ID	AAS03687/c		
XX	AAS03687	standard; DNA; 3093 BP.	
AC	AAS03687;		
XX			
DT	29-AUG-2001	(first entry)	
XX			
DE	Rhesus gene locus: RHD gene deletion in Rh negative haplotypes.		
XX			
KW	Rhesus box: RHD positive; sequence length polymorphism; SSP; RHD; SMP1;		
KW	RHCP: Rh negative; blood group typing; blood transfusion; antigen C;		
KM	haemolytic disease of the newborn; chromosome 1 p34.1-p36; ds.		
XX			
OS	Homo sapiens.		
XX			
FH	Key	Location/Qualifiers	
FT	primer_bind	32..54	
FT		/*tag= a	
FT	primer_bind	/note= "Binding site of primer rez"	
FT		Complement (3034..3054)	
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FT		/note= "Binding site of primer rnd31"	
XX			
PN	W0200132702-A2.		
XX			
PD	10-MAY-2001.		
XX			
PF	31-OCT-2000; 2000WO-EP10745.		
XX			
PR	02-NOV-1999; 99EP-0121686.		
PR	31-MAY-2000; 2000EP-0111696.		
XX			
PA	(DRKB-) DRK BLUTSPENDEDIENST BADEN WUERTTEMBERG.		
XX			

PI Flegel WA, Wagner FF;
XX
DR WPI: 2001-291052/30.
XX
PT New nucleic acid molecular structure, useful for detection of common
PT RHD positive haplotypes in D-negative individuals, comprises RHD, SMP1
PT and RHCE genes -
XX
PS Example 10; Fig 5; 135pp; English.
XX
CC The sequence represents the coding sequence of Rhesus gene locus:
CC RHD gene deletion in Rh negative haplotypes. The Rhesus genes
CC locus comprises the RHD, SMP1 and RHCE (all undefined) genes and/or the
CC Rhesus box(es), preferably the hybrid Rhesus box, the upstream Rhesus box
CC and/or the downstream Rhesus box. The RHD and RHCE genes are located at
CC chromosome 1 p34.1-p36. Rhesus box flanks the breakpoint region of the
CC RHD deletion in the common RHD negative haplotypes. The sequence has
CC been used to design primers which are useful for: (1) the specific
CC detection of the common RHD positive haplotypes in D-negative
CC individuals; (2) blood group typing; (3) determining whether a patient
CC can be transfused with Rhd negative blood and whether blood is suitable
CC for transfusion to patients who should not be exposed to antigen C; (4)
CC assessing the risk of a Rhd negative mother of conceiving or carrying an
CC Rhd positive foetus. Anti-D antibodies are useful for treating pregnant
CC women who are Rhesus D negative, where the foetus is not homozygous for
CC the RHD gene to treat or prevent haemolytic disease of the newborn.
XX
XX Sequence 3093 BP; 891 A; 754 C; 619 G; 829 T; 0 other:
50

Query Match 32.8%; Score 452; DB 22; Length 3093;
Best Local Similarity 47.4%; Pred. NO. 1.1e-103;
Matches 485; Conservative 212; Mismatches 282; Indels 45; Gaps 5;

Qy 390 rgargaratbsnaarcarcarwsnathcargargtnacntgygntlnylnaargcnt 449
Db 1338 AGGAGAAATTTCTTAGCAAGCAAAAGCATTCAGAGGTGACTGGCTGTAAAGACATT 1279
Qy 450 ywsntlytngngargngargcaayaarwsnsgaraayncayncngayayagntat 509
Db 1278 CAGTTTAAAGAGGAGGAGCAAGCATTAAGGTTGGAATTTGGACGCTGCAAGTGTAT 1219
Qy 510 haaraaraaraa--ycntlywsngargnaartlyaarlytngcngcngarathtyath 567
Db 1218 AGAANAAGAAATTTCCATTTCTGAGGAGAAATTCAGAGTGTGCGAGAAATTTGCATG 1159
Qy 568 tgyaaygarlythaayglnaaycncargaaayagngaraayaalhtwsntgacntgy 627
Db 1158 AGTAAC-AGGAGCCAAATGCTAATTCCCAAGACAAATGGGGAATGTCTCCAGGCGATGT 1100
Qy 628 carmgwswncarcarcarsnatlhaarsnyng--cntgmgncnmngmaartgft 686
Db 1099 CAGAGGCTCTTATGTCACACCCCTCCCATCAGGTCAGAGGTATCAGGAAAAATGTTT 1040
Qy 687 ytygynacngcngcngnswntlytgytlygtnarcnmngnngaytngtncntlygt 746
Db 1039 TTGTTGGCCAGGCGCGGGGTCTCATGCTGTGACGCTAGGAGCTTGCTGCTGCAT 980
Qy 747 nccngtna-----aywsngcngtngcwsng 772
Db 979 CCCAGCCACTCCCAACCATGACTGAGGAGGAGCAAGGTAGAGCTTGCGCTTGCTTCGG 920
Qy 773 argngcwmwncncnarcncntgsgaryncnswngngtngcngtngngcnaara 832
Db 919 GGAGTGCAGAGCCCCAAGCCTTGACAGCTTCCATGTGTGTGAGCTGCGAATGCACAA 860
Qy 833 arwsnmatlbgargtngtngcncnathngntlycaraaarlthaygnaaycnc 892
Db 859 AGTCAAGATTTGGGTTGGAAACCTTGCGCTAGATTAAAGAGATGTGCGGAATGCTT 800
Qy 893 ggaatcmmgncaraarttycngtngcngtngcngtngcngtngcngtngcngtngcng 952
Db 799 GGATGCCAGTCAGAAATTTGCTGTCAGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGG 740

Qy 953 tngtncaraargnaaytngntggarcncncncaymgntncngngcncnc 1012
Db 739 CAGTCAGAGAGGAAATTTGGGTGAGAGACCCACACACAGTCCCTACTGGGGACACAC 680
Qy 1013 snwsnmgncngtngnmgngnswncncnswnsnmgntncaraargngmgngnswncng 1072
Db 679 CTAGTGAGAGCTGTGAGAGAGAGGTCTCC-----AGACCCAGAAATGTGATCACCG 627
Qy 1073 aywsntncarcaytngcncngaraarwsnncngayncartgyarcngtngnaargcng 1132
Db 626 ACAGCTTGACCGTGTACTGTGAAAAAGCTGCAGACACTCAATGACACGCCCATGAAAGCAG 567
Qy 1133 cngnatlgarwsngtngcncntayaaraengtngtngcngtngcngtngcnaarngtngna 1192
Db 566 CTGAGAGGAGGAGCTTACCTCCACCAAGGTACAGGCGCAGAGGTGCCCAAGACATGGGAA 507
Qy 1193 thlaytngtncaytgycaayaytngaytngmncaygngtngnaarmngaycaelytgy 1252
Db 506 CCCACCCCTTCATCTGCTGCTGACGTGATGTGAGATGTGAGATCAAGAGATCATTTTG 447
Qy 1253 gngcngtngmngtlyaytgygcncngntlymgnaactayatyggngcngtngcngtnt 1312
Db 446 GAGCTTTAAGATTTGACTGCCCCACTGATTTTGGACTCTCATGAGGCGCTGTAGCCTTT 387
Qy 1313 gtytygncartlytlycncntlygngnagcngtntlyacncartgytlnlaytncayt 1372
Db 386 GTTTGACCAATTTTATCCATTTGGAATGGCTGTATTATCCCAATGCTGTATCCCCATT 327
Qy 1373 gyat 1376
Db 326 GTAT 323

RESULT 3
AAS03689/C
ID AAS03689 standard; DNA; 9236 BP.
XX
AC AAS03689:
XX
DT 29-AUG-2001 (first entry)
XX
DE Rhesus gene locus: upstream Rhesus box of D-positives.
XX
KW Rhesus box: RHD positive; sequence length polymorphism: SSP; RHD; SMP1;
KW RHCE; Rh negative; blood group typing; blood transfusion; antigen C;
KW haemolytic disease of the newborn; chromosome 1 p34.1-p36; ds.
XX
OS Homo sapiens.
XX
PN W0200132702-A2.
XX
PD 10-MAY-2001.
XX
PF 31-OCT-2000; 2000WO-EPI0745.
XX
PR 02-NOV-1999; 99EP-0121686.
PR 31-MAY-2000; 2000EP-0111696.
XX
PA (DRKB-) DRK BLUTSPENDEDIENST BADEN WUERTEMBERG.
XX
PI Flegel WA, Wagner FF;
XX
DR WPI: 2001-291052/30.
XX
XX New nucleic acid molecular structure, useful for detection of common
XX PT RHD positive haplotypes in D-negative individuals, comprises RHD, SMP1
XX PT and RHCE genes -
XX
XX Disclosure: Fig 9; 135pp; English.
XX
CC The sequence represents the coding sequence of Rhesus gene locus:
CC upstream Rhesus box of D positives. The Rhesus genes locus

CC comprises the RHD, SMPI and RHCE (all undefined) genes and/or the
CC Rhesus box(es), preferably the hybrid Rhesus box, the upstream Rhesus box
CC and/or the downstream Rhesus box. The RHD and RHCE genes are located at
CC chromosome 1 p34.1-p36. Rhesus box flanks the breakpoint region of the
CC RHD deletion in the common RHD negative haplotypes. The sequence has
CC been used to design primers which are useful for: (1) the specific
CC detection of the common RHD positive haplotypes in D-negative
CC individuals; (2) blood group typing; (3) determining whether a patient
CC can be transfused with Rhd negative blood and whether blood is suitable
CC for transfusion to patients who should not be exposed to antigen C; (4)
CC assessing the risk of a Rhd negative mother of conceiving or carrying an
CC Rhd positive foetus. Anti-D antibodies are useful for treating pregnant
CC women who are Rhesus D negative, where the foetus is not homozygous for
CC the RHD gene to treat or prevent haemolytic disease of the newborn.

SQ Sequence 9236 BP; 2467 A; 2319 C; 2000 G; 2450 T; 0 other;

Query Match 32.8%; Score 452; DB 22; Length 9236;
Best Local Similarity 47.4%; Pred. No. 2.7e-103;
Matches 485; Conservative 212; Mismatches 282; Indels 45; Gaps 5;

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QY 390 rgararathwnaarcarcarsnaathcargatnactaggtgtnlytnaargcntt 449
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Db 6014 AGAGAGAAATTCTTAAGCAGCAAGCATTCAAGAGTGACTTGCGTCTTAAAGCATT 5955
QY 450 ywsntlyathmgngarngarcayaaarwsnsgaraaylncayccngayaaygnat 509
  :||:||||: ||:||||: ||:||||: ||:||||: ||:||||: ||:||||: ||:||||:
Db 5954 CAGTTTATTAAGAGAGACAGACATTAAGTTGCGAAATTTGCAAGCCTGCAATGTGAT 5895
QY 510 haaraaraara--ycenctlywsngarngnaartlyaaalytngcngaratngyath 567
  :||:||||: ||:||||: ||:||||: ||:||||: ||:||||: ||:||||: ||:||||:
Db 5894 AAAAAAGAAAAATTCCTATTTCTGAGGAAATTCACCTGCGTGCACAAATTTGCAAG 5835
QY 568 tgyaaagargarytlnaaylncaycncargayaaygngarayaayltnsngacntly 627
  :||:||||: ||:||||: ||:||||: ||:||||: ||:||||: ||:||||: ||:||||:
Db 5834 AGTAAC-AGGAGCCAAATCTTAATTCACAGCAATGGGAAATGTCTCCAGGGCATGT 5776
QY 628 carmgwnswnsncarcarsnaathaarwsnylnc-ntlgmngncmngmnaartggt 686
  :||:||||: ||:||||: ||:||||: ||:||||: ||:||||: ||:||||: ||:||||:
Db 5775 CAGAGGTCTTTATGGCAACCCCTCCATCAGAGTCCAGAGATACAGAAAAAATGCTT 5716
QY 687 ytgynagncngncngncngnwsnylntgytgytlnccmngngayltnctnctgyt 746
  :||:||||: ||:||||: ||:||||: ||:||||: ||:||||: ||:||||: ||:||||:
Db 5715 TGTGTGGCCAGGCCGCGGGTCTCATGTGTGTCAGCCTAGGAGCTTGTGCTCGCAT 5656
QY 747 nccngtlna-----aywsngcngncngnwsng 772
  :||:||||: ||:||||: ||:||||: ||:||||: ||:||||: ||:||||: ||:||||:
Db 5655 CCCAGCCACTCCCAACCATGACTGACGGAGCCAGGCTAGAGCTTGAGCTTCGG 5596
QY 773 argngcwnsncnaarcnctlgcarylncnwsngnglncarcngltnngncnaara 832
  :||:||||: ||:||||: ||:||||: ||:||||: ||:||||: ||:||||: ||:||||:
Db 5595 GGAGTCCAGGCCCAAGCCTTGACAGCTTCATGTGCTTGAGACTCGGAGGCACAGA 5536
QY 833 arwsnmgnathgargtntggarcncncaatlmngntlyaraarathaygnaaycnc 892
  :||:||||: ||:||||: ||:||||: ||:||||: ||:||||: ||:||||: ||:||||:
Db 5535 AGTCAAGAAATGGGTTGGAAACCTTCCTAGATTAAAGAGATGTCCGGAATAATGCTT 5476
QY 893 ggaatgcmmngnaraartlycngltnngltnngltnngmwnsngnagncnwsngmng 952
  :||:||||: ||:||||: ||:||||: ||:||||: ||:||||: ||:||||: ||:||||:
Db 5475 GGATGCCCGCTAGAAATTTGCTGCAGGAGCAAGGCCCTCATGAGAAATCTCTGCGAGG 5416
QY 953 lnglncaraargnaaygltngltnngltnngarcncncaaymngltnncnwsngngncncw 1012
  :||:||||: ||:||||: ||:||||: ||:||||: ||:||||: ||:||||: ||:||||:
Db 5415 CAGTGCAGAGGAAATGTGGGGTGCAGAGACCCACACATCCCTACTGGGGCACCAC 5356
QY 1013 snwsnmngcngltnmgngnwsncnccnwsnwsnmngnylncaraargmngnwsnag 1072
  :||:||||: ||:||||: ||:||||: ||:||||: ||:||||: ||:||||: ||:||||:
Db 5355 CTAGTGCAGCTGTGAGAAAGAGTCTCC-----AGACCCCAAGATGTAGATCCACCG 5303
QY 1073 aywsnylncarcayltnccngararwsnagayncnarygarcngltnngnagng 1132
  :||:||||: ||:||||: ||:||||: ||:||||: ||:||||: ||:||||: ||:||||:
Db 5302 ACAGCTTGCAACCGTGTACTGAAAAGCTGCAGACACTCAATGCCCATGAAAGCAG 5243
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QY 1133 cngnatggarwsnlnccntlayaaraacngltnngcngaryltnaanaacngltnnga 1192
  :||:||||: ||:||||: ||:||||: ||:||||: ||:||||: ||:||||: ||:||||:
Db 5242 CTGAGAGGAGGCTGTACCTCTCAAAAGGTACAGGGGACAGCTGCCCAAGACCATGGGA 5183
QY 1193 thtalytlncaaygycaaygyltnngaygtmngcnyvngngltnaarnngayccytlg 1252
  :||:||||: ||:||||: ||:||||: ||:||||: ||:||||: ||:||||: ||:||||:
Db 5182 CCCAGCCCTCTGCTGCTGACCTGGATGTGATGTGAGTCAAGAGATCATTTTGG 5123
QY 1253 gngcnyltnmgntlyayltygcnaacngnlytmgnacntlayatggngcngltnctnt 1312
  :||:||||: ||:||||: ||:||||: ||:||||: ||:||||: ||:||||: ||:||||:
Db 5122 GAGCTTTAAGATTTGACTGCCCTGATTTTGCACTCTCATGGGCTGTAGCCTCTTT 5063
QY 1313 gyltygncartlytlycncntlyggnacngcngltnlycnaacartgyylnalytncayt 1372
  :||:||||: ||:||||: ||:||||: ||:||||: ||:||||: ||:||||: ||:||||:
Db 5062 GTTTTGACCAATTTATTCATTTGGAAATGCTGTATTATCCCAATGCCCTGATCCCAT 5003
QY 1373 gytat 1376
  :||:
Db 5002 GTAT 4999
```

RESULT 4

AA03690/C
ID AA03690 standard; DNA; 9236 BP.

AA03690;

29-AUG-2001 (first entry)

Rhesus gene locus: downstream Rhesus box of D-positives.

Rhesus box; RHD positive; sequence length polymorphism; SSP; RHD; SMPI;
RHCE; Rh negative; blood group typing; blood transfusion; antigen C;
haemolytic disease of the newborn; chromosome 1 p34.1-p36; ds.

Homo sapiens.

WO200132702-A2.

10-MAY-2001.

31-OCT-2000; 2000WO-EP10745.

02-NOV-1999; 99BP-0121686.

31-MAY-2000; 2000BP-0111696.

(DRKB-) DRK BLUTSPENDEDIENST BADEN WÜRTTEMBERG.

Fliegel WA, Wagner FF;

WPI; 2001-291052/30.

New nucleic acid molecular structure, useful for detection of common
RHD positive haplotypes in D-negative individuals, comprises RHD, SMPI
and RHCE genes -

Disclosure: Fig 10; 135pp; English.

The sequence represents the coding sequence of Rhesus gene locus:
downstream Rhesus box of D positives. The Rhesus genes locus
comprises the RHD, SMPI and RHCE (all undefined) genes and/or the
Rhesus box(es), preferably the hybrid Rhesus box, the upstream Rhesus box
and/or the downstream Rhesus box. The RHD and RHCE genes are located at
chromosome 1 p34.1-p36. Rhesus box flanks the breakpoint region of the
RHD deletion in the common RHD negative haplotypes. The sequence has
been used to design primers which are useful for: (1) the specific
detection of the common RHD positive haplotypes in D-negative
individuals; (2) blood group typing; (3) determining whether a patient
can be transfused with Rhd negative blood and whether blood is suitable
for transfusion to patients who should not be exposed to antigen C; (4)
assessing the risk of a Rhd negative mother of conceiving or carrying an
Rhd positive foetus. Anti-D antibodies are useful for treating pregnant

[illegible]

ID	AA005134	standard; cDNA; 3203 BP.
XX		
AC	AA005134;	
XX		
DT	17-JUL-2001	(first entry)
XX		
DE	Human secreted protein-encoding gene 14 cDNA clone H15BF60, SEQ ID NO:24.	
XX		
KW	Human; secreted protein; proliferative disorder; cancer; tumour;	
KW	fetal abnormality; developmental abnormality; haematopoietic disorder;	
KW	immune system disorder; AIDS; autoimmune disease; rheumatoid arthritis;	
KW	inflammation; allergy; neurological disorder; Alzheimer's disease;	
KW	Parkinson's disease; cognitive disorder; schizophrenia; asthma;	
KW	skin disorders; psoriasis; sepsis; diabetes; atherosclerosis;	
KW	cardiovascular disorder; angogenic disorder; kidney disorder;	
KW	gastrointestinal disorder; pregnancy-related disorder;	
KW	endocrine disorder; infection; wound healing; vulnery;	
KW	cell culture; chemotaxis; food additive; gene therapy;	
KW	binding partner identification; ss.	
XX		
OS	Homo sapiens.	
XX		
FT	Key	Location/Qualifiers
FT	CDS	153..356
FT		/*tag= a
FT	sig_peptide	/product= "Human secreted protein"
FT		153..215
FT		/*tag= b
FT	mat_peptide	216..353
FT		/*tag= C
FT		/product= "Human mature secreted protein"
XX		
PN	WO200134769-A2.	
XX		
PD	17-MAY-2001.	
XX		
PF	01-NOV-2000; 2000WC-US30040.	
XX		
PR	05-NOV-1999; 99US-0163580.	
PR	30-JUN-2000; 2000US-0215130.	
XX		
PA	(HUMA-) HUMAN GENOME SCI INC.	
XX		
PI	Ruben SM, Komatsoulis GA, Wei P, Baker KP, Fiscella M;	
XX		
DR	WPI: 2001-308781/32.	
DR	P-PSDB: AAE01245.	
XX		
PT	New isolated nucleic acid molecule encoding a human secreted protein is	
PT	used in preventing, treating or ameliorating a medical condition -	
XX		
PS	Claim 1; Page 408-409; 519pp; English.	
XX		
CC	AA005121-AA005203 represent cDNAs corresponding to 24 human secreted	
CC	protein genes, and AA001232-AA001311 represent the proteins they encode.	
CC	AA001312-AA001340 represent human secreted protein variants or fragments	
CC	The secreted proteins and their genes are useful for preventing,	
CC	treating or ameliorating medical conditions, e.g., by protein or gene	
CC	therapy. Pathological conditions can be diagnosed by determining the	
CC	amount of the new protein in a sample or by determining the presence of	
CC	mutations in the new genes. Specific uses are described for each of the	
CC	24 genes, based on the tissues in which they are most highly expressed,	
CC	and include developing products for the diagnosis or treatment of	
CC	proliferative disorders, cancer, tumours, foetal and developmental	
CC	abnormalities, haematopoietic disorders, diseases of the immune system,	
CC	AIDS, autoimmune diseases (e.g., rheumatoid arthritis), inflammation,	
CC	allergies, neurological disorders (e.g., Alzheimer's disease,	
CC	Parkinson's disease), cognitive disorders, schizophrenia, asthma,	
CC	skin disorders (e.g., psoriasis), sepsis, diabetes, atherosclerosis,	
CC	cardiovascular disorders, angiogenic disorders, kidney disorders,	
CC	gastrointestinal disorders, pregnancy-related disorders, endocrine	
CC	disorders, and infections. The proteins can also be used to aid wound	
CC	healing and epithelial cell proliferation, to prevent skin aging due to	

Oy	688	tcygnacngcngcngcngcngnylnctyctgctncaacnmgagaytngtncctggttn	747
Db	3779	tcgtggccagagcccaagggtccccaatgctgtgtgcagcccaaggagcttggtccctgcac	3838
Oy	748	ccnglnaabyngs-----cnglncngwsngary	775
Db	3839	ccagctgctccagcatctgtctaaaggccgcgaggtacagctctgcgcctgtgttcaarg	3898
Oy	776	gngcwnscncaarccnltgcarytncnwsngngtngarcngtngngcnaaragw	835
Db	3899	gtgcagagcccaaaccttgccagcttccatctgtgtgtctgagcctgcagtgcatgaagct	3958
Oy	836	smngnathrgcngtngggarcnccnccnaltmngtltayaraaralthaygnaayccttga	895
Db	3959	caagaatttgaggttggttggaacctccatctagataltcagaagatgtatgtgaatcaccttga	4018
Oy	896	tgccmngncararttctgacngtngtngtngtngwsnmsnctgmgngnacsngcmmngtng	955
Db	4019	tgctccagcgaagaaagcttgcttccaggggcagagaccttcatgaggaacccctgtctaggcag	4078
Oy	956	lncaraaaygntngngtngtngggarcnccnccncaymngtncnwsngngcncnwsnw	1015
Db	4079	tg-tgaagggaatgttggtgtgtgagccccaacagaaatccctactgtgggacaccta	4137
Oy	1016	smngngcngtngmgngnwsnccnccnwsnwsnmgnylncaraargmgngnwsncaangyw	1075
Db	4138	gtcgagcgtgtgcagaagaagagccacgcttcccttcagaccacgaattgtatctccagca	4197
Oy	1076	snynccarcaygtncnccnccnccnccnccnccnccnccnccnccnccnccnccnccnccn	1134
Db	4198	gctgtgacgtgcacccctgcgaaagccacagaaacccataacgcagcccgagaaagca	4257
Oy	1135	gngatgawrsngtncnccnccnccnccnccnccnccnccnccnccnccnccnccnccnccn	1194
Db	4258	ggagtggtggagctataccctgtgagccaaagggcgagcgtgcacaaagcttaaggaaoc	4317
Oy	1195	taytngtncaytgcycagaytngaygtngmgncaygngtlnaarmngaycaytlyggn	1254
Db	4318	tacccttgcatcatctgtgtgacccgtgagtcagacatgagtcagagagatatttga	4377
Oy	1255	gcnymngtcttgcaytgcycnccnccnccnccnccnccnccnccnccnccnccnccnccn	1313
Db	4378	acgttataattctacgtgcctcgtgattcttggaacttgacagggagctgtgaaagccttg	4437
Oy	1314	yttcngcarytctyccncttgcngcngcngtntcncnccnccnccnccnccnccnccnccn	1373
Db	4438	ttgtgtggacacttccctccatttggaatggtctgtattacccttaccgtatacccatgt	4497
Oy	1374	yatg 1377	
Db	4498	tatg 4501	
RESULT 8			
AAS44505 standard; DNA; 22509 BP.			
AAS44505:			
AAS44505:			
18-DEC-2001 (first entry)			
Human LEKTI DNA clone C1978SKB_94F21 contig 11, SPINK5 exons 1-4.			
Human: SPINK5; lympho-epithelial Kazal-type related inhibitor; LEKTI; ds			
serine protease inhibitor; atopic disease; Netherton's syndrome; asthma;			
eczema; hayfever; antihistaminic; antiallergic; antiinflammatory;			
dermatological; PCR primer; sequencing primer; gene therapy.			
Homo sapiens.			
W0200164747-A1.			
07-SEP-2001.			

PF	02-MAR-2001	2001MO-GB00897.	
XX			
XX	02-MAR-2000	2000GB-0005098.	
PR	03-MAR-2000	2000GB-0005229.	
XX			
PA	(ISIS-) ISIS INNOVATION LTD.		
PI	Hovnanian A, Chavanas S, Cookson W, Moffat M, Walley A;		
PI	MDJ; 2001-582149/65.		
XX			
DR			
PT	Determining susceptibility to atopic disease or carrier status of		
PT	Netherton's syndrome in humans by identifying variants of or mutations		
PT	in SPINK5, a gene encoding lympho-epithelial Kazal-type related		
PT	inhibitor		
XX			
PS	Disclosure; Page 88-94; 123pp; English.		
CC			
CC	Sequences AAS44359-AAS44514 represent the SPINK5 gene, contigs and		
CC	fragments of a SPINK5 clone, sequencing primers and PCR primers for		
CC	SPINK5. SPINK5 encodes lympho-epithelial Kazal-type related inhibitor		
CC	(LEKTI), a serine protease inhibitor. Susceptibility or predisposition to		
CC	an atopic disease in a human subject can be detected by screening the		
CC	genome for one or more polymorphic variants of SPINK5 gene and/or		
CC	expression of a variant LEKTI protein in a tissue. Carrier status of a		
CC	subject or development of Netherton's syndrome is diagnosed by screening		
CC	for the presence of loss-of-function mutations in the SPINK5 gene. An		
CC	expression vector comprising a nucleic acid encoding a serine protease		
CC	inhibitor or its functional fragment can be used in screening for		
CC	compounds with potential pharmacological activity by determining the		
CC	serine protease activity of a protein previously identified as a ligand		
CC	of the LEKTI protein. The atopic diseases include Netherton's Syndrome,		
CC	asthma, eczema and hayfever.		
XX			
SO	Sequence 22509 BP; 6725 A; 4291 C; 4284 G; 7209 T; 0 other;		
Query Match	31.0%; Score 427.2; DB 22; Length 22509;		
Best Local Similarity	47.1%; Pred. No. 1,1e-96;		
Matches	472; Conservative 207; Mismatches 303; Indels 21; Gaps 7;		
QY	390 rgatgarathwnsaarcarcarwsnathcargaratnactgggtgntynynaargcnt 449		
DB	14187 agaagaattcttaagcagcagcagcatcaagaagataacttgggtgcttaagcatt 14246		
QY	450 ywsnttlyatlmngnargcngarcayaaarwsnwsngarayaaytlncaocngayaaygtat 509		
DB	14247 cagctcttlaaggaagacagacataaaagtttggaaatttgcagctgcagctgatt 14306		
QY	510 haataaata--rraayccnttlywsngatgnaarttlyaaaytngcngcngarathtgyath 567		
DB	14307 aaaaaaagaataatccattcttcgaggaagaattccaagcaagctgtcataattgtcata 14356		
QY	568 tgyaaygarayrlnaaygltlnaayccncaarayaayngngaayathwsnltgacntgy 627		
DB	14367 agtgaagaaggcctgtatgttaatccccaagaacattgg- gaaatgtctccaagaagat 14425		
QY	628 carmgwnswncarcarwsnathaarwsyntgcntgymngcmmgmgnaar---tgg 684		
DB	14426 cagaagaccttctgbcagccctccatcatatcccaagaggttlaaggaataaaatgg 14485		
QY	685 tctygygnacngcngcmgnwsytnfytgytlncaocngmmngaytltngtncntgy 744		
DB	14486 tttccttagccaggtcctaaggtctccctgcgtgtgcagccctcgltgaaagacattca 14545		
QY	745 gtncncg-----tlnaaycngcngtngcmwsngaragngcwnsccncaarccntg 794		
DB	14546 gtgcctgaaggagccatgctgtgycgtatgcttccgaggtgcgaagccttaagccctg 14605		
QY	795 gcaaytncmwsngngtngarcngcngtngngcnaaraarwsmmgnathargntltgga 854		
DB	14606 acacacttccaaatgttcttgagcctgttgaagtcacagaagccaagaacttgagttgaaa 14665		

Db 14842 gccctcatccctccagagaaccagatatagacatccacagtagcttgaccatagtctga 14901
QY 1095 raarrwnaacngayacncaatgycarcnngtnaargcngcngnatggarwnnglncnta 1154
Db 14902 gaaagccacaacaactcagctgagccagcctgtgaagcgagcgaggagcggtlaccctg 14961
QY 1155 yaaraacngtgtngcngarytlnacnaaraacngtngnathlaytlnaytlnacayga 1214
Db 14962 caaaagccacagagagtgagagtgccccaagaccatgaggaaaccacctctgacacatga 15021
QY 1215 ytlngaygtlmngncaygngtnaarmngaycaytlygngcnytlmngntlygaytgyc 1274
Db 15022 cctgagatagagacatgattcaaaatatacttttgaaactttaagatttgactgcct 15081
QY 1275 naacngntlymnaacntaytggncngt-nccnyntlytlygncatlytlycct 1333
Db 15082 tccctgaatttcagactgcatgcatctgtagcccttlttgagcacaatctcccat 15141
QY 1334 tygnaacngcngtlytynacnartgytlnaytlnaytlnaytlyat 1376
Db 15142 ttggaacagctgtatttgacccaatgctgtaccccatgtat 15184

RESULT 10
ID AAL03634
XX AAL03634 standard; DNA; 6063 BP.
AC AAL03634:
DT 21-NOV-2001 (first entry)
XX
DE Human reproductive system related antigen DNA SEQ ID NO: 6322.
XX
KW Human; reproductive system related antigen; reproductive system disorder;
XX cancer; gene therapy; ds.
OS Homo sapiens.
XX
PN MO200155320-A2.
PD 02-AUG-2001.
XX
PF 17-JAN-2001; 2001MO-US01339.
XX
PR 31-JAN-2000; 2000US-0179065.
PR 04-FEB-2000; 2000US-0180628.
PR 24-FEB-2000; 2000US-0184664.
PR 02-MAR-2000; 2000US-0186350.
PR 16-MAR-2000; 2000US-0189874.
PR 17-MAR-2000; 2000US-0190076.
PR 18-APR-2000; 2000US-0198123.
PR 19-MAY-2000; 2000US-0205515.
PR 07-JUN-2000; 2000US-0209467.
PR 28-JUN-2000; 2000US-0214886.
PR 30-JUN-2000; 2000US-0215135.
PR 07-JUL-2000; 2000US-0216647.
PR 07-JUL-2000; 2000US-0216880.
PR 11-JUL-2000; 2000US-0217487.
PR 11-JUL-2000; 2000US-0217496.
PR 14-JUL-2000; 2000US-0218290.
PR 26-JUL-2000; 2000US-0220963.
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PR 14-AUG-2000; 2000US-0224518.
PR 14-AUG-2000; 2000US-0224519.
PR 14-AUG-2000; 2000US-0224519.
PR 14-AUG-2000; 2000US-0225213.
PR 14-AUG-2000; 2000US-0225214.
PR 14-AUG-2000; 2000US-0225266.
PR 14-AUG-2000; 2000US-0225267.
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PR 14-AUG-2000; 2000US-0225270.
PR 14-AUG-2000; 2000US-0225477.
PR 14-AUG-2000; 2000US-0225477.
PR 14-AUG-2000; 2000US-0225757.
PR 14-AUG-2000; 2000US-0225758.

PR 14-AUG-2000; 2000US-0225759.
PR 18-AUG-2000; 2000US-0226279.
PR 22-AUG-2000; 2000US-0226681.
PR 22-AUG-2000; 2000US-0226868.
PR 22-AUG-2000; 2000US-0227182.
PR 23-AUG-2000; 2000US-0227009.
PR 30-AUG-2000; 2000US-0228924.
PR 01-SEP-2000; 2000US-0229287.
PR 01-SEP-2000; 2000US-0229343.
PR 01-SEP-2000; 2000US-0229344.
PR 01-SEP-2000; 2000US-0229345.
PR 05-SEP-2000; 2000US-0229509.
PR 05-SEP-2000; 2000US-0229513.
PR 06-SEP-2000; 2000US-0230437.
PR 06-SEP-2000; 2000US-0230438.
PR 08-SEP-2000; 2000US-0231242.
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PR 08-SEP-2000; 2000US-0231414.
PR 08-SEP-2000; 2000US-0231414.
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PR 08-SEP-2000; 2000US-0232081.
PR 12-SEP-2000; 2000US-0231968.
PR 14-SEP-2000; 2000US-0232397.
PR 14-SEP-2000; 2000US-0232398.
PR 14-SEP-2000; 2000US-0232399.
PR 14-SEP-2000; 2000US-0232400.
PR 14-SEP-2000; 2000US-0232401.
PR 14-SEP-2000; 2000US-0233063.
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PR 21-SEP-2000; 2000US-0234223.
PR 21-SEP-2000; 2000US-0234274.
PR 25-SEP-2000; 2000US-0234997.
PR 25-SEP-2000; 2000US-0234998.
PR 26-SEP-2000; 2000US-0235484.
PR 27-SEP-2000; 2000US-0235834.
PR 27-SEP-2000; 2000US-0235836.
PR 29-SEP-2000; 2000US-0236327.
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PR 02-OCT-2000; 2000US-0236802.
PR 02-OCT-2000; 2000US-0237037.
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PR 02-OCT-2000; 2000US-0237039.
PR 02-OCT-2000; 2000US-0237040.
PR 13-OCT-2000; 2000US-0239937.
PR 20-OCT-2000; 2000US-0240960.
PR 20-OCT-2000; 2000US-0241221.
PR 20-OCT-2000; 2000US-0241785.
PR 20-OCT-2000; 2000US-0241786.
PR 20-OCT-2000; 2000US-0241787.
PR 20-OCT-2000; 2000US-0241808.
PR 20-OCT-2000; 2000US-0241809.
PR 20-OCT-2000; 2000US-0241826.
PR 01-NOV-2000; 2000US-0244617.
PR 08-NOV-2000; 2000US-0246474.
PR 08-NOV-2000; 2000US-0246475.
PR 08-NOV-2000; 2000US-0246476.
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PR 08-NOV-2000; 2000US-0246523.
PR 08-NOV-2000; 2000US-0246524.
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PR 08-NOV-2000; 2000US-0246526.
PR 08-NOV-2000; 2000US-0246527.
PR 08-NOV-2000; 2000US-0246528.
PR 08-NOV-2000; 2000US-0246532.
PR 08-NOV-2000; 2000US-0246609.
PR 08-NOV-2000; 2000US-0246610.

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PR	17-NOV-2000;	2000US-0249224.	
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PR	17-NOV-2000;	2000US-0249226.	
PR	17-NOV-2000;	2000US-0249227.	
PR	17-NOV-2000;	2000US-0249228.	
PR	17-NOV-2000;	2000US-0249229.	
PR	17-NOV-2000;	2000US-0249230.	
PR	01-DEC-2000;	2000US-0250391.	
PR	05-DEC-2000;	2000US-0251030.	
PR	05-DEC-2000;	2000US-0251988.	
PR	05-DEC-2000;	2000US-0256719.	
PR	06-DEC-2000;	2000US-0251479.	
PR	08-DEC-2000;	2000US-0251856.	
PR	08-DEC-2000;	2000US-0251866.	
PR	08-DEC-2000;	2000US-0251869.	
PR	08-DEC-2000;	2000US-0251989.	
PR	08-DEC-2000;	2000US-0251990.	
PR	11-DEC-2000;	2000US-0254097.	
PR	05-JAN-2001;	2001US-0259678.	
XX			
PA	(HUMA-)	HUMAN GENOME SCI INC.	
XX			
PI	Rosen CA, Barash SC, Ruben SM;		
XX			
DR	WPI; 2001-465570/50.		
XX			
PT	Isolated nucleic acid molecule encoding a reproductive system antigen		
XX	Is used in preventing, treating or ameliorating a medical condition -		
XX			
PS	Disclosure; SEQ ID NO 6322; 1297bp + Sequence Listing; English.		
CC			
CC	The present invention provides the protein and coding sequences of a		
CC	number of human reproductive system related antigens. These can be used		
CC	in the prevention and treatment of reproductive system disorders,		
CC	including cancer. The present sequence is a genomic sequence encoding a		
CC	protein of the invention.		
XX			
SO	Sequence 6063 BP; 1656 A; 1163 C; 1262 G; 1982 T; 0 other;		
Query Match	31.0%; Score 426.4; DB 22; Length 6063;		
Best Local Similarity	46.4%; Pred. No. 5.9e-97;		
Matches	475; Conservative 210; Mismatches 299; Indels 40; Gaps 6;		
OY	331 gargarathswnaarcarcawrnatbthargargtnaentggtnlytnlynaargcntly 450		
Db	3461 gaaggaattcttaagcagcaagaacatccaagagtgatcttggtcgtgttaagaacatt 3540		
OY	451 wntlyahmgngagcngarcarayaaarwmsngaraayyncaycngayaaaytnath 510		
Db	3541 cattttaaaggaac--agcataaaaatttgaaaatttgacgcacgttatgcagca 3598		
OY	511 aaraaraaayccctlytwngargngaaattlyaaayngcngcngarathtgyathgy 570		
Db	3599 gaagaggaagaacactcttttttgaggaagaattccaagcgcgcgcgaatttgataagt 3658		
OY	571 aaygaraglynaaygtnaayccnargayaayagngaraayathwnttgacntgycar 630		

Db	3659	tacaggagacgtgaattgtttaactctccaagacacatgagggaataatgtctccagggcgtgtcac	3718
Oy	631	mgunswnsncarwarwnathaarwsnlytngcrltgmgncmmgmgnaar---	Egatty 687
Db	3719	agctctcatatgacgacccctcccatcaacacccctgaagactagggaggaataatggttt	3778
Oy	668	lgygnaacngnccngnswsnlytlgytlnearcmmngayytlnglncnltgyltn	747
Db	3779	lgttggccgaagcccgaggtcccatgtgctgtgtgaagcttagggacttggctgcctgtcatc	3838
Oy	748	ccngtnaaywng-----cnglncmwsngatg	775
Db	3839	ccagctgtctccagccattgtctaaaggccgaagtgtaacgtctgcctgtggttccagag	3898
Oy	776	gngcmwsnccnaarccnltgcaarylncmwsngnltngarcnngtngnngcnnaarw	835
Db	3899	gttcaagccccaacaccttggaagcttccatctgtgtgttgagccttggaagltgcatggaaat	3958
Oy	836	snmgnaatgharglnttggarccnccnaalmnglttycaaraathlayggaanaayc	895
Db	3959	caagaattgaggttgaggaaacctcatctagatttcaagaagatgtatgtaacatcgtga	4018
Oy	886	ttccmmnncatnaattlycngltngngtngngnswsnltgmgnaecmwsngcmngtng	955
Db	4019	tgctcaagccaaaagtltgtctccagggaagcagctcatcttgagaaacctctgttaagngc	4078
Oy	956	lncraaraaggaagyltngnltggarccnccmcamngnltncmwsngnngnccmwsnw	1015
Db	4079	tg-tgaagggaaatgtgtgggtlttgagagccccaacagaaatccctactgtggacaccta	4137
Oy	1016	snmgncnglmgngmgnwsnccnccmwsnsmngnlylncraaraagmgmgwnacngayw	1075
Db	4138	gttgaagctgtgcaagaagccacacgtctctcaagaccacgaatgtgtagatccacgaca	4197
Oy	1076	snylncarcaaygttncngaraarawnaengayacnaartgycaarccngtnaarngcn	1134
Db	4198	gcttgtaccgtgcaccttggaataagccacaacctaaccgacccagccgttgaaagcagcca	4257
Oy	1135	gnaatlgarwsngnccntlayaarcngtltngcngarylnaarcnarrcngltngnath	1194
Db	4258	ggagggggagactatacccttggaagccacaggggcagagctgtgccaaagactaaggaaac	4317
Oy	1195	taytlnylncayltycagayytlngaygtlmgncaygtlmaazmgnaaycaaytlytgn	1254
Db	4318	taccctctgcacatctgtgaacctgagtgtagacacagggatcgaaggaagatacttttga	4377
Oy	1255	gcnylmngnttygaytyccmaecngnltymnancntayaltgngncnglt-nccnylnb	1313
Db	4378	acgtataataatgagctctgcctgcgtgatttttgactatgcatggtggtcttaacgcttg	4437
Oy	1314	ytytgnaactlytgcnttlytngnaecngnlttyaacnactgtytnttaytlncaayt	1373
Db	4438	lttgggggaactctcccatcttgaaatggtctgattataccattaccgttatacccatgt	4497
Oy	1374	yatg 1377	
Db	4498	tatg 4501	
RESULT 11			
ID	AAH13678	AAH13678 standard; cDNA; 1736 BP.	
XX	AAH13678;		
XX	26-JUN-2001 (first entry)		
XX	Human cDNA sequence seq ID NO:10539.		
XX	Human; primer; detection; diagnosis; antisense therapy; gene therapy; ss		
XX	Homo sapiens.		
XX			

PM WO200162969-A2.
XX
PD 30-AUG-2001.
XX
PF 20-FEB-2001; 2001WO-US05358.
XX
PR 22-FEB-2000; 2000US-183756P.
PR 20-OCT-2000; 2000US-0692414.
PR 24-JAN-2001; 2001US-0768184.
XX
PA (PEKE) PE CORP NY.
PI Kalush F, Cassel MJ, Hwang SS, Winn-Deen ES;
XX WPI: 2002-041152/05.
DR P-PSDB; AAG68251.
XX
PT Novel variant of estrogen receptor alpha polypeptide useful for
PT determining the biological activity of a protein for high throughput
PT screening and for raising antibodies that elicit an immune response in
PT host .
PS Example: Fig 1 page 1-93; 333pp; English.
XX
CC The present invention describes an isolated peptide (I) consisting of an
CC amino acid sequence selected from: (a) the amino acid sequence of a
CC variant of the estrogen receptor alpha (ESR-alpha) protein in AAG68251;
CC or (b) a fragment comprising at least 10 contiguous amino acids of the
CC protein in AAG68251. (I) has cytostatic, osteopathic, cardiant and
CC vasotropic activities, and can be used in gene therapy and vaccine
CC production. (I) is useful for identifying an agent that binds to (I), by
CC contacting (I) with an agent and assaying the contacted mixture to
CC determine whether a complex is formed with the agent bound to the
CC peptide. A polynucleotide (II), encoding (I), is useful in the
CC development of diagnostics and therapies for diseases and disorders
CC mediated/modulated by an estrogen receptor (ER). (II) is also useful in
CC gene therapy for treating cancer, osteoporosis and cardiovascular
CC diseases. The human ESR-alpha gene is located on chromosome 6. The
CC present sequence represents the human ESR-alpha gene, which is given in
CC the exemplification of the present invention.
XX
SQ Sequence 465237 BP; 133988 A; 89578 C; 93946 G; 147721 T; 4 other;

Query Match 30.6%; Score 421; DB 24; Length 465237;
Best Local Smlarity 46.5%; Pred. No. 5e-94;
Matches 477; Conservative 213; Mismatches 292; Indels 43; Gaps 7;

QY 392 argarathmsnaarcaraarwathcargartnactgggttynynaargcntlyw 451
DB 438788 aagaatattctaagcagcaaaacatccaaggctgacttggtgtgttaaaagcatctt 438847
QY 452 snlyatlmngnargcnarcarayaaarwswngnaraaytlnacaycngayaatnatha 511
DB 438848 gttttaaaagagaac--agctlaaagtlcagaatatttgagcctgtagtcagtag 438905
QY 512 aaraaraaaycc-nlytwsngargnagarttyaarytngcngnargathtyaltgyt 570
DB 438906 gaagaagaaacccattttttgaggaatccaagctgcgcagaaattgtcataagt 438965
QY 571 aaygararytnaaygtlnaaycncargayaayngnaraaytlnatwsntgscntgycar 630
DB 438966 aacaagagaccaaattgttaattcccaagaacaatlggggaaatcgtccagacatgcat 439025
QY 631 mgnwswnsncarcatwatharwtnlyngnttgmgncmngmng---naartggtt 686
DB 439026 aggttctcttgagcagccctcccatcacagaccggaagccttagagagaaaaaacagt 439085
QY 687 ytygynacngcngcngnwnsnyntlytygtynacnngmngngaytlnatncntgyt 746
DB 439086 tttgtgacagtcacagaggtcccatgtgtgtgacagcctagaaatgtgtccctgcat 439145
QY 747 ncnngtnaaywngcngtngc-----nwnngatrgngcnws 782

DB 439146 ctgagctgtccagcatattgtccaaagggtcgtgagttaccacggttccaaggttgcag 439205
QY 783 nccnaarcentgyccarctnccnswngngtngarccngtngngcnaaarwsmngnat 842
DB 439206 ccccaaaccttgagcagcttctcatgtggtgttgagccttgltgtaacagaagttagaat 439265
QY 843 hgarctntggagccnccnathmngntlycaaraarcthaaygnaayccttgatgcmmg 902
DB 439266 tgaagttggaaacctccatctatattccaagaagaatacgttgnaatgtcctgataccag 439325
QY 903 ncaaraattgcnngtngngtngtngnswntgngnccnswngcngmngtngtccaara 962
DB 439326 gcaaacattgtcgcagaggtggtggccccaagagggcctctgtaggacatgaggaa 439385
QY 963 rgnnaaytngnttgtagccnccncaaymgngtncnswngngcngcnnswsmngnc 1022
DB 439386 gggaaatgtgggtgtggaaccccccaacagagttcccaactgggacagcctagtgagc 439445
QY 1023 ngtnmgngnswncnccnswnsnmngntncaraarngmngnswnaengaywsnytnca 1082
DB 439446 tylaagagagagaccacgtctctccagaccgcagagatagtagatccatgacatgtgca 439505
QY 1083 rcaygtncngarrraarwsmnacngayacncarctgyarccngtlnaargcngcngna---- 1138
DB 439506 ccatgtccttggaagaagcacagacacctaacgcagccttgaagaagcatgaggttg 439565
QY 1139 -----tggarwsngtncntayaaarccngtngtngcngartnacaaraongtngna 1192
DB 439566 aggtgtgtgtgtctatacccttaagccacagagggcagagctgtcccaagctaggaa 439625
QY 1193 thtaytlnacaytgycaaygtngaygtlmgncayggngtnaarmngaycaaytgy 1252
DB 439626 cctacccttgcatcagacatgacgtcgtgtagcatctcagtcacaaagaaat-atttg 439684
QY 1253 gngcnytmngntlyaytygycnaongntlymgnaactlayatggngcngt-nccnytn 1311
DB 439685 aagcttgaatctgacgtgcctgtgtgatttagactgtgtggccctgaacccctt 439744
QY 1312 tgytngncarttytccnttlygnaongcngntlyacncaatgytntaytlnca 1371
DB 439745 tgtttgccaattctcccatlttgtagctgtgtagtattaccatgtccaaaccgcat 439804
QY 1372 tgyat 1376
DB 439805 tgtat 439809

RESULT 13
AAS91984
ID AAS91984 standard; cDNA; 1485 BP.
XX
AC AAS91984;
XX
DT 13-FEB-2002 (first entry)
XX
DE DNA encoding novel human diagnostic protein #27788.
XX
KW Human; chromosome mapping; gene mapping; gene therapy; forensic;
KW food supplement; medical imaging; diagnostic; genetic disorder; ss.
XX
OS Homo sapiens.
XX
PN WO200175067-A2.
XX
PD 11-OCT-2001.
XX
PF 30-MAR-2001; 2001WO-US08631.
XX
PR 31-MAR-2000; 2000US-0540217.
PR 23-AUG-2000; 2000US-0649167.
XX
PA (HYSE-) HYSEQ INC.

PE 01-NOV-2000; 2000MO-US30040.
XX 05-NOV-1999; 990S-0163580.
PR 30-JUN-2000; 2000US-0215130.
XX
PA (HUMA-) HUMAN GENOME SCI INC.
XX
XX Ruben SM, Komatsoulis GA, Wei P, Baker KP, Fisceella M;
PI WPI; 2001-308781/32.
XX P-PSDB; AAE01289.
DR
XX
PT New isolated nucleic acid molecule encoding a human secreted protein is
PT used in preventing, treating or ameliorating a medical condition -
XX
XX
PS Claim 1; Page 442; 519pp; English.
XX
CC AAD05121-AA005203 represent cDNAs corresponding to 24 human secreted
CC protein genes, and AAE01232-AAE01311 represent the proteins they encode.
CC AAE01312-AAE01340 represent human secreted protein variants or fragments.
CC The secreted proteins and their genes are useful for preventing,
CC treating or ameliorating medical conditions, e.g., by protein or gene
CC therapy. Pathological conditions can be diagnosed by determining the
CC amount of the new protein in a sample or by determining the presence of
CC mutations in the new genes. Specific uses are described for each of the
CC 24 genes, based on the tissues in which they are most highly expressed,
CC and include developing products for the diagnosis or treatment of
CC proliferative disorders, cancer, tumours, foetal and developmental
CC abnormalities, haematopoietic disorders, diseases of the immune system,
CC AIDS, autoimmune diseases (e.g., rheumatoid arthritis), inflammation,
CC allergies, neurological disorders (e.g., Alzheimer's disease,
CC Parkinson's disease), cognitive disorders, schizophrenia, asthma,
CC skin disorders (e.g., psoriasis), sepsis, diabetes, atherosclerosis,
CC cardiovascular disorders, angiogenic disorders, kidney disorders,
CC gastrointestinal disorders, pregnancy-related disorders, endocrine
CC disorders, and infections. The proteins can also be used to aid wound
CC healing and epithelial cell proliferation, to prevent skin aging due to
CC sunburn, to maintain organs before transplantation, for supporting cell
CC culture of primary tissues, to regenerate tissues, to identify their
CC cognate ligands or binding partners, and in chemotaxis, and can be used
CC as a food additive or preservative to modify storage properties.
CC Antibodies specific for a protein of the invention can be used in
CC alleviating symptoms associated with the disorders mentioned above, and
CC in diagnostic immunoassays e.g., radioimmunoassay or enzyme linked
CC immunosorbent assay (ELISA). The present sequence represents a human
CC secreted protein-encoding cDNA of the invention.
XX
XX
SQ Sequence 1278 BP; 344 A; 264 C; 358 G; 299 T; 13 other;

Query Match 29.7%; Score 408.8; DB 22; Length 1278;
Best Local Similarity 45.8%; Pred. No. 4.8e-93;
Matches 458; Conservative 205; Mismatches 298; Indels 40; Gaps 6;

QY 391 gargarathwsnaarcacarsnathcargartnacntggtntlytynaargcntly 450
DB 280 gaagaataattcctcaagcaagaagcattcaagaagtgattgggtctgtttaaagcattt 339
QY 451 wntlyatlmngnargcngarcayarswngaraaytlncaycngayaaygtatnath 510
DB 340 cattttaaagggaac--agcataaaatttgaaatttcagcagcagtgatgcagaa 397
QY 511 aaraaraaaycncntlywsngargnarttyaarytngcngcngarathtgtyathgy 570
DB 398 gaagagaagaacctgttttttgagagaattcaagctggcgggaatttgcaatagt 457
QY 571 aaygargarlythaaygtlnaaycncargayaaygngnaraaaythwsntgagcngtycar 630
DB 458 tacagggagcgtgaatgttaattccaagaacaatggggaataatgtctccagggcatgtcac 517
QY 631 mgnwsnscarcaswathaarwnytnngntlymgngncmngmgnaar--tggty 687
DB 518 aggtcttcatgtgcagccctcccatcacagaccctgaagactagagaaaaatggttt 577

QY 688 tgytgagacngnngcngnswntlytgytncarcnmngnayaenlytncntlygt 747
DB 578 tgytgagcagccagccaggtcccatctgctgtgacagctagagacttgcctgtcatc 637
QY 748 ccngtlnaayngcngtngc-----nwsngarg 775
DB 638 ccagctgtctccagccatttgcataaaggccagaggtacagctctgcgccatgttcaagagg 697
QY 776 gngcwnsncnaarcnltgycarytncnwsngngtngarcnngtngngnmaaraarw 835
DB 698 gtgcaagcccaaaccttgcagcttcacatgtgtgttgacgttcgacaggtcagatg 757
QY 836 smngnatlhargntltggargcnccnaltmgntlycaraarathlaygnaaycmtgga 895
DB 758 caagaaltgaggttgyggaacctccatcagatcagaaagatgcatgaatccaccgga 817
QY 896 tgcnmngncaraarttlycngtngngtngngtngnswnttggmngnacnwsngcmmngtng 955
DB 818 tgcctcagcaaaaagtctgtctccagggcagagccttcattgagaaacctctcagggcag 877
QY 956 tncaraargnaaygtngngtnggarcncncaymngntncnwsngngcncnsw 1015
DB 878 tg-tgaagggaatgtgggttggagcccyacacagaatccctactggggccacctta 936
QY 1016 smngngcngtngmngnswncnswncnswsmngntncaraargmngnswnacngayw 1075
DB 937 gtgagacgtgtcggaagaagccagccagctgtccagaccctcagaaatggtgtgacccagca 996
QY 1076 snytncrcaygtncngaraarsnagayacnartlycarcngtlnaargcng-cn 1134
DB 997 gcttgcacgycgacaccttgaaagacacagaccctlaagcagcccgltgaagcagca 1056
QY 1135 ggnatggrswngtncnttaraaengtngtngcngarytlnaanaaengtngnath 1194
DB 1057 ggaatgggactataccctgtgaagcccaagggcagagctgcgcgaagaactaaaggaa 1116
QY 1195 taytlnycaytgcaygaaytngaygtlmngnagngtlnaarmngaycaytlyggn 1254
DB 1117 taactctgcacatctngnaccctgnatgttgagatgagatgagagagagatcatttigna 1176
QY 1255 gonytmgntlygaatgycncaacngnttymgnacntayatlvgngcngtncnlyntly 1314
DB 1177 acgnataatttgactgcctgcgtgatttgagacttgcattgaggtcgtgnaacgccttlt 1236
QY 1315 tlyg-gnarttlytncntlyngnagcngcngtnttycnc 1354
DB 1237 gtgtgagacttctcccatctggaatggtcgttattacc 1277

RESULT 15
AAS87262/C
ID AAS87262 standard; cDNA: 1580 BP.
XX
XX AAS87262;
XX
XX 13-FEB-2002 (first entry)
XX
XX DNA encoding novel human diagnostic protein #23066.
XX
XX Human; chromosome mapping; gene mapping; gene therapy; forensic;
XX food supplement; medical imaging; diagnostic; genetic disorder; ss.
XX
XX Homo sapiens.
XX
XX WO200175067-A2.
XX
XX 11-OCT-2001.
XX
XX 30-MAR-2001; 2001WO-US08631.
XX
XX 31-MAR-2000; 2000US-0540217.
XX 23-AUG-2000; 2000US-0649167.
PR

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XX (HYSE-) HYSEQ INC.
PA
XX Drmanac RT, Liu C, Tang YT;
XX
XX MPI: 2001-639362/73.
XX P-PSDB; ABG3075.
XX
XX New isolated polynucleotide and encoded polypeptides, useful in
XX diagnostics, forensics, gene mapping, identification of mutations
XX responsible for genetic disorders or other traits and to assess
XX biodiversity.
XX
XX Claim 1: SEQ ID NO 23066; 103pp; English.
XX
XX The invention relates to isolated polynucleotide (I) and
XX polypeptide (II) sequences. (I) is useful as hybridisation probes,
XX polymerase chain reaction (PCR) primers, oligomers, and for chromosome
XX and gene mapping, and in recombinant production of (II). The
XX polynucleotides are also used in diagnostics as expressed sequence tags
XX for identifying expressed genes. (I) is useful in gene therapy techniques
XX to restore normal activity of (II) or to treat disease states involving
XX (II). (II) is useful for generating antibodies against it, detecting or
XX quantitating a polypeptide in tissue, as molecular weight markers and as
XX a food supplement. (II) and its binding partners are useful in medical
XX imaging of sites expressing (II). (I) and (II) are useful for treating
XX disorders involving aberrant protein expression or biological activity.
XX The polypeptide and polynucleotide sequences have applications in
XX diagnostics, forensics, gene mapping, identification of mutations
XX responsible for genetic disorders or other traits to assess biodiversity
XX and to produce other types of data and products dependent on DNA and
XX amino acid sequences. AAS64197-AAS94564 represent novel human
XX diagnostic coding sequences of the invention.
XX Note: The sequence data for this patent did not appear in the printed
XX specification, but was obtained in electronic format directly from WIPO
XX at ftp.wipo.int/pub/published.pat_sequences.
XX
XX Sequence 1580 BP; 357 A; 412 C; 328 G; 483 T; 0 other;
XX
Query Match 29.1%; Score 400.4; DB 23; Length 1580;
Best Local Similarity 42.4%; Pred. No. 7.6e-91;
Matches 466; Conservative 215; Mismatches 383; Indels 34; Gaps 4;
QY 307 aaytayaywmsnltgyngarytncaycaytygvaarytlnaayahtygtnatg 366
DB 1107 AATTCGATACGACAGACAGCTGTGCTACTAGTTCAGTTCAGCCCGGAGCACTCA 1048
QY 367 mgnaaacathytngcnaaaygaargarathwsnaarcarwarasnathcargatn 426
DB 1047 GCATCCCCCACTTCTTAGAGGACAGAGTTCAGCCACTGAGATTGAGCAATAC 988
QY 427 acntgggtngtynatnaagcnclywsntlyatlmgngargcngarcayaaarwmsngar 486
DB 987 AGATGTGGTTTGAATTTGAATTTATTTAAAGGAGACAGCATTAATAAATTGGA 928
QY 487 aaytyncaycngayaaycgnathaaaraaaraaycnclywsngargnaartlyaar 546
DB 927 AATCTGACGATGACAGTGAATTAAGAAAAGAAACCCATTCTTGCGGAGAAATTCAG 868
QY 547 ytingcngcngarahtcayathcyaaargarytlnaaytlnaaycncargayaaygn 606
DB 867 CTGCTTGACAAATTTGATTAATTAACAGGACCAATGTTAATCCAAAGACACGGG 808
QY 607 garaayathwntgagacnlycarmgnwmsnscarcarwsnathaaarwntylngcngt 666
DB 807 AAAAATGTTTACAGGACGACGACCTTATGCAAGCCCTCCCATGCGAGCCAG 748
QY 667 mgnccmngmngnaa---rtgtytygngacngcngcngnwnytnlytygtncar 723
DB 747 AGGCTTAGGAGAAATATGTTTGTGGCTGGCCAGGACCTTGCTGCTTTGTGACG 688
QY 724 ccmgngaytytngtncntgylngcngtlnaaywsng----- 760

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DB 687 TCTCAGACTTGTGCTCCCTGCATCCATCTGTGCTAAAGGGCCAAATACAGCTGAA 628
QY 761 -cngtngcngwngargngcngwscnccnaarccntbgcarytncngsnngngngarccn 819
DB 627 ACCATTGCTTCAAGAGGTGTAAAGCCCAAGCATTTGAGAGCTGATGTTGTGACCT 568
QY 820 gtnngcnaaaraarwmsnmgnaathcargtntgggarcncncnathmgntlycaraarath 879
DB 567 GTGGGTACACAGAAAGTCACAACTGAGGTTGGGAACTCTGCTGATTTACAGAGATG 508
QY 880 taygnaaycnclygaltgcccmgncaraartlycngtngngngtngngwmsnlygmgm 939
DB 507 TATGGAACACCTGATATATCAGAGAGGTGCTGACAGGGGTAGAGCCCTCATGAGAG 448
QY 940 acwmsngcmngtngtncaraargnaaytngtngtngtngtngtngtngtngtngtng 999
DB 447 ACCTCTGCTAGGCAATGACAGAAAGGAAATGTGGGCTTGAGAGCTCCACACAGATCCC 388
QY 1000 wsnngcngcngwmsnmgngcngtngmngmgnwscncncwmsnmgngtncaraar 1059
DB 387 ACTGG-----GACTGTGAGCTGTGAGAAAGGGCCACCTCTCCAGACCCAGAAAT 334
QY 1060 gmgmgnwscnagaywnylncarcaytncngararaarwscnagayancartlycar 1119
DB 333 GATAGACCAACTGACAGCTTGATGCTGTGACCTGGAAGAAACCAACCAACTCAATGCAAT 274
QY 1120 cngtlnaargcngcngatgargwngtncntlayaarecngtngtngcngarytncn 1179
DB 273 CCTGTGAACACACGAGAGGGAGGAGCTGTACCTGCAAAACCCAGAGGGCAGAGCTTCC 214
QY 1180 aaraecngtngnahtbaytncaytncaytycaygarytngaytngtngcngngtncar 1239
DB 213 AAGGCTGTAAACCCACCTTTACATACAGCTGTCTTGATGTGACCTGAGATCAAA 154
QY 1240 mgnaycaytlygngcngtngmngntlygalytgcnaacngntlymgnaactayatlgn 1299
DB 153 GAGATCATTTTGGAGATTTTAAAGATTTGGCTGCCCCACTGATTTAGAGATGATGGG 94
QY 1300 cngt-ngcnytnlytlytngcncartlytlycnclytngcngcngtngtlycncartg 1358
DB 93 CCTGTACCCCTTGTGTTTGGCCAAATTTCTCCATCTGGAATGAGTGTGTTATCCAAATG 34
QY 1359 ytnlaytncaytyat 1376
DB 33 CCCGTACCCCATTTATAT 16

```

Search completed: July 8, 2002, 16:20:51
Job time: 12424 sec

...

GenCore version 4.5
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OM nucleic - nucleic search, using sw model

Run on: July 8, 2002, 12:57:56 ; Search time 104.62 Seconds
(without alignments)
3233.008 Million cell updates/sec

Title: US-09-997-610-3

Perfect score: 1377

Sequence: 1 aethgtnatnccngtnt.....gytntatyytncatgyatg 1377

Scoring table:

Gapop 10.0 , Gapext 1.0 .

Searched: 383533 seqs, 122816752 residues

Total number of hits satisfying chosen parameters: 767066

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Listing first 45 summaries

Database :

Issued_Patents_NA: *
1: /cgn2_6/ptodata/2/1na/5A.COMB.seq: *
2: /cgn2_6/ptodata/2/1na/5B.COMB.seq: *
3: /cgn2_6/ptodata/2/1na/6A.COMB.seq: *
4: /cgn2_6/ptodata/2/1na/6B.COMB.seq: *
5: /cgn2_6/ptodata/2/1na/PCTRUS.COMB.seq: *
6: /cgn2_6/ptodata/2/1na/backfiles1.seq: *

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	299.2	21.7	49136	US-09-422-869-1	Sequence 1, Appl
2	177.6	12.9	14855	US-08-687-080-59	Sequence 59, Appl
3	164.6	12.0	620	US-08-991-789A-29	Sequence 29, Appl
4	164.6	12.0	620	US-09-062-451-29	Sequence 29, Appl
5	140.6	10.2	152331	US-09-128-155-16	Sequence 16, Appl
6	135.8	9.9	14855	US-08-687-080-59	Sequence 59, Appl
7	67	4.9	7218	US-08-232-463-14	Sequence 14, Appl
8	59	4.3	2067	US-09-106-194-11	Sequence 11, Appl
9	51.8	3.8	1313	US-08-463-911-6	Sequence 6, Appl
10	51.8	3.7	1155	US-09-140-804-9	Sequence 9, Appl
11	50.6	3.7	2082	US-09-053-866-3	Sequence 3, Appl
12	48.8	3.5	2082	US-09-440-325A-2	Sequence 2, Appl
13	48.6	3.5	289	US-09-351-414-3	Sequence 3, Appl
14	48.6	3.5	289	US-09-007-005-17	Sequence 17, Appl
15	48.6	3.5	289	US-09-244-796-17	Sequence 17, Appl
16	48.6	3.5	1107	US-09-188-930-217	Sequence 217, App
17	48.6	3.5	1276	US-08-463-911-1	Sequence 1, Appl
18	48.2	3.5	2277	US-08-676-967-2	Sequence 2, Appl
19	48.2	3.5	2277	US-08-676-974-2	Sequence 2, Appl
20	48.2	3.5	2277	US-08-676-967-2	Sequence 2, Appl
21	48	3.5	2277	US-08-676-967-2	Sequence 2, Appl
22	48	3.5	2277	US-09-098-487-2	Sequence 2, Appl
23	48	3.5	2277	US-09-098-487-2	Sequence 2, Appl
24	44.4	3.2	1212	US-09-092-770-18	Sequence 18, Appl
25	44.4	3.2	1212	US-09-222-851-18	Sequence 18, Appl
26	43.4	3.2	5181	US-08-257-073-10	Sequence 10, Appl
27	43	3.1	1215	US-09-092-770-8	Sequence 8, Appl

C	28	43	3.1	1215	4	US-09-222-851-8	Sequence 8, Appl
C	29	42.4	3.1	1015	3	US-09-188-930-30	Sequence 30, Appl
C	30	42	3.1	2067	4	US-09-106-194-11	Sequence 11, Appl
C	31	41.8	3.0	1001	3	US-09-188-930-218	Sequence 218, App
C	32	41.8	3.0	2265	3	US-09-369-618-3	Sequence 3, Appl
C	33	41.8	3.0	2265	3	US-09-369-617-3	Sequence 3, Appl
C	34	41.6	3.0	510	2	US-08-934-959-7	Sequence 7, Appl
C	35	41	3.0	597	4	US-09-528-760A-3	Sequence 3, Appl
C	36	40.8	3.0	723	3	US-08-911-423-5	Sequence 5, Appl
C	37	40.8	3.0	1659	1	US-08-231-729B-2	Sequence 2, Appl
C	38	40.6	2.9	2949	4	US-09-412-554A-3	Sequence 3, Appl
C	39	40.2	2.9	500	3	US-09-141-000-2	Sequence 2, Appl
C	40	40.2	2.9	1659	1	US-08-231-729B-1	Sequence 1, Appl
C	41	40	2.9	2088	4	US-09-351-414-3	Sequence 3, Appl
C	42	39.8	2.9	2265	3	US-09-369-618-3	Sequence 3, Appl
C	43	39.8	2.9	2265	3	US-09-369-617-3	Sequence 3, Appl
C	44	39.6	2.9	4379	1	US-08-592-214A-17	Sequence 17, Appl
C	45	39.6	2.9	4379	3	US-09-149-976-17	Sequence 17, Appl

ALIGNMENTS

RESULT 1
US-09-422-869-1
: Sequence 1, Application US/09422869
: Patent No. 6235481
: GENERAL INFORMATION:
: APPLICANT: POLONSKY, KENNETH S.
: APPLICANT: HORIKAWA, YUKIO
: APPLICANT: ODA, NAOHISA
: APPLICANT: COX, NANCY J.
: APPLICANT: SREENAN, SEANUS
: APPLICANT: ZHOU, YUN-PING
: APPLICANT: OTANI, KENICHI
: APPLICANT: HANIS, CRAIG L.
: APPLICANT: BELL, GRAEME I.
: TITLE OF INVENTION: METHODS OF TREATMENT OF TYPE 2 DIABETES
: FILE REFERENCE: ARCD:307
: CURRENT APPLICATION NUMBER: US/09/422,869
: EARLIER FILING DATE: 1999-10-21
: EARLIER APPLICATION NUMBER: 60/134,175
: EARLIER FILING DATE: 1999-05-13
: NUMBER OF SEQ ID NOS: 30
: SOFTWARE: PatentIn Ver. 2.0
: SEQ ID NO 1
: LENGTH: 49136
: TYPE: DNA
: ORGANISM: Human
: US-09-422-869-1

Query Match 21.7%; Score 299.2; DB 4; Length 49136;
Best Local Similarity 41.1%; Pred. No. 1.1e-72;
Matches 388; Conservative 186; Mismatches 307; Indels 62; Gaps 5;

OY	444	rgcnttysnttyahmgngaragcngarcayaaarsnsgaraaylncayccngayaa	503
OY	504	yttnatharaaraaraayccntlywsngargnaarttyaarylncngcngarathg	563
DB	47384	ggaactccatttaaaaggaagcagagcataaaagtttaaaattgcatcccgta	47443
OY	47444	tgagataaaaaaaactcatcttcggagaggaatcaagccagctgcagaattg	47503
OY	564	yathtyaayagaryltnaayccnaryayaaygngaraayathwsntgac	623
DB	47504	catagtaactagagccacatgtaataagcatagacaatgaggaatgtccaagc	47563
OY	624	ntgyarngnswncarcawsnatthaarwsntgntgngncmmgnaa	680
DB	47564	atgtcagaggtcttcacagcaaccacccatcaagcctgagagccttaagagga	47623
OY	681	rtggttygyggnacngncngnwsnylntgtytgcrtncarccmngaylntgnc	740

Db	47624	atgctgtcgtcgtcgtccagagcctctgctgtgtcgtcgtcagcttggtgc	47683
Qy	741	ntggttncngtnaaywsgnngtgcmsnsgarngcwnsnccnaarcttgcaat	800
Db	47664	ccaaatcc-----cagcagctgctaaagggccaaatgataagcttaagctt	47731
Qy	801	ncnwnsgngtngarcnccngtngngcnaaraarwsmgnathgarltngtgarcncc	860
Db	47732	tgctcagaagtgcaagc-----	47755
Qy	861	natlmngnttlycaaraathaygnaaycngtgcacgcmngncararctlycngtng	920
Db	47752	caagccttggtgctcttaactgtgtgttgctggcctgcagatacagaagcttgctgcact	47811
Qy	921	ngtngnswntbngmnaacnswngcmgngtngtlnaaraargnaaygtngtngtgg	980
Db	47812	ggtgtaaccctcaatgataacccctgcagagcgagctgtagaagtgataagtgvggttga	47871
Qy	981	rcnccncaymngtncnwnsgngncc-----cwnswsmngnccngtmgmgnws	1034
Db	47872	gccccccacaacaatcccaactgvggcactgctactgtactgtgaactgtgagaagaag	47933
Qy	1035	ncnccnswsmngnythncaraargmngmwnsacngaywntynhncarctaytncnnga	1094
Db	47932	gccaccatcccccagaccaccaagtgatagataccactgtactgtgaaccaagcactgg	47991
Qy	1095	raarwnacngayaacnartgycarcnngtlnaargcngcngnatgarnswngtncnna	1154
Db	47992	aaagccacagacactcaaccacagcctgtgaaagcagctggaagggagcgtacccg	48055
Qy	1155	yaarccngtngtngcngarythnaaraacngtngnathaytynhncaytgcayga	1214
Db	48052	caaaacaacagagcagagctgcccagaaggtcatggagagccacccttgcatagaagctga	48111
Qy	1215	yytngaaytlnmcnagcngnhtnaarmngayaaytlytngcngnyhnmgttgaaytgc	1274
Db	48112	cttgaaatgtgaacatgagatcaagaagatcatcttggaagcttaagaatcttgacggcc	48171
Qy	1275	nacnganttymnactatayattgagccngt-ncnynltgtytlyggncaarttytgcnt	1333
Db	48172	acctgattctggaacttgatgcagtgggcctgtgccccttcatttlttgccaattatcccat	48233
Qy	1334	tygganacngcngtnttynacnartgtythtatytncaaytgc	1376
Db	48232	ctggaatggatattatcaaccatgctcgtaccatctcat	48274

```

APPLICATION NUMBER: US 08/592,126
FILING DATE: 26-JAN-1996
ATTORNEY/AGENT INFORMATION:
NAME: Sholtz, Charles K.
REGISTRATION NUMBER: 38,615
REFERENCE/DOCKET NUMBER: 4600-0111.30
TELECOMMUNICATION INFORMATION:
TELEPHONE: (415) 324-0880
TELEFAX: (415) 324-0960
INFORMATION FOR SEQ ID NO: 59:
SEQUENCE CHARACTERISTICS:
LENGTH: 14855 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
HYDROTHERMAL: NO
ANTI-SENSE: NO
ORIGINAL SOURCE:
INDIVIDUAL ISOLATE: 5' END OF INTRON 2 OF RAD50 GENOMIC
INDIVIDUAL ISOLATE: SEQUENCE
US-08-687-080-59

Query Match      12.9% Score 177.6; DB 2; Length 14855;
Best Local Similarity 44.4%; Pred. NO. 7.7e-39;
Matches 221; Conservative 90; Mismatches 172; Indels 15; Gaps 4.

QY   890  cntgatcgccmgnncgaarcttlygngtngtggtgmgswnsntgmgnaacnwgncgm  949
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Db   14854  CCTGATGCCCGACAGCAGATGTTCCTGCACGGGGTGGGCCCTTAGGAACCTCTGCTA  14795

QY   950  gngltngtcaraazrgnaaygtngtnltygarccnccncaymngtlcnccwnsgngcnc  1009
       : : : : | : | : | : | : | : | : | : | : | : | : | : | : | : |
Db   14794  GGCGAAATATGGAAGGAATGTGGGTGGAACCCACC - AGAGTTCTCATGAGGGGAC  14737

QY   1010  cnwswsmgmngcngtlnmgmgmwscnccnccnswsmmgnyl-----ncarraar  1059
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Db   14736  TGCCATAGTGAGACTGTGTGAGAAACACAGCCACTGTCTCCAGACTGTATGATCCCGAGAT  14677

QY   1060  gnmgnwnsnacgawswyntlncarcaygtncngaraarwsnacgayaencartgycar  1119
       : : : : | : | : | : | : | : | : | : | : | : | : | : | : | : |
Db   14676  AATAATATCACACGACAGCTGTGCATCTGTCACTCGAAMAACTGCAGGCACTATACACACAG  14617

QY   1120  ccngtlnaargcngcngnatlgayarsvngtncntayaaarcngltngtngcngarytnaacn  1179
       : : : : | : | : | : | : | : | : | : | : | : | : | : | : | : |
Db   14616  CCGTGTGAATAAACAGCCAGAGAGAGGCGCTATACCTCCCAAAGCC - AGAAGTGAGAGCGCCC  14559

QY   1180  aaareongtngnatltxaytuytlncaiygycaysayytngaygtlmnpagaygnptnaar  1239
       : : : : | : | : | : | : | : | : | : | : | : | : | : | : | : |
Db   14558  AAGGCCATGGAAGCCCCACTCTTGCAATCAGAGTAAGCTGATGTGAGACATGAGAGTCANA  14459

QY   1240  mngnacycatlytgngcngnytlmtngatltlyaglygcnaacngncltymgnaentlayabggn  1299
       : : : : | : | : | : | : | : | : | : | : | : | : | : | : | : |
Db   14498  GGAGATCAATTCGTGAGCCTTAGATATACACCTCCCOACTGAAATTTCGGACTTGACAGGGG  14439

QY   1300  ccngf-nccynfntlytlygncatltlytcentltlygnaengcngtltlacnearg  1358
       : : : : | : | : | : | : | : | : | : | : | : | : | : | : | : |
Db   14438  CCGTGAAGCCCTCTTCTTTGGCCAATTTCTCCCAATTGGATGGAGGCTGATATTGGCCCAATG  14379

QY   1359  yyintaytlncaiyatlyat  1376
       : : : : | : | : | : | : | : | : | : | : | : | : | : | : | : |
Db   14378  CCTGTATCCCATTTGTAT  14361

RESULT      3
US-08-991-789A-29
Sequence 29, Application US/08991789A
Patent No. 6225054
GENERAL INFORMATION:
APPLICANT: Frudakis, Tony N.
Smith, John M.
Reed, Steven G.
```



```
Db 183 TACTAGAAAGTACAGAGAAATGTGGTTGGAGCCCCCAACAGAAATCCCTCTAG 242
Qy 1005 ngnccnswsmngnngntlmngmgnwscnccnswsmngnylncaaraagmg 1064
Db 243 AACACCTGCTAATGAACCTGTGAGAAATGGCCACTGTCAATCCAGACACCAATGTAG 302
Qy 1065 nwnacnngaynynlncarcayg-lncngaraarwnacnngayncartlycarcc-- 1121
Db 303 ACCCACCAAAAACCTATATGCTATATAAAACCTACAGACACTCAATGACGAGCCCC 362
Qy 1122 -ngnaarngcnngnatgatgarwsnnglncnlaayaraarngntngcngarlna 1180
Db 363 ATGAAAAAAAACAGAGAGAGAGACTGTCCTCAATGCCACCGAGAGCAACTGCCCC 422
Qy 1181 araengnngnathlaytynlncaycayagaylncayg-lmngcayagngtnaar 1239
Db 423 AGGCGATGGAAGACACAGCTGTATATCATGTACCTGAGATGTGAGCAATGAAATCMA 482
Qy 1240 mngnagayltyngnngnltmngntltygaytycnaengnngntlmngnactay 1295
Db 483 NGAATGCTTTTAACTCCACGCTTAAATGACTGCCCTATTAATTCNCACTTANAT 542
Qy 1296 g-gnccngnccnngntlytyltyngncaartlytyngnngnngnngntlync 1354
Db 543 CCNCGCTGTGACCTCTTGTCTTGCCATTCCTTTTGGAAAGGCTMTTTTTC 602
Qy 1355 artylntaylnc 1369
Db 603 CATGCTGTNCCCTC 617

RESULT 5
US-09-128-155-16
; Sequence 16, Application us/09128155
; Patent No. 6117654
; GENERAL INFORMATION:
; APPLICANT: Pan, Yang
; TITLE OF INVENTION: NOVEL MOLECULES OF TANGO-77 RELATED PROTEIN FAMILY
; FILE REFERENCE: 09404/052001
; CURRENT APPLICATION NUMBER: US/09/128,155
; EARLIER FILING DATE: 1998-08-03
; EARLIER APPLICATION NUMBER: US 60/091,650
; EARLIER FILING DATE: 1998-07-02
; EARLIER APPLICATION NUMBER: US 60/054,646
; NUMBER OF SEQ ID NOS: 18
; SOFTWARE: FASTSEQ for Windows Version 3.0
; SEQ ID NO 16
; LENGTH: 152331
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)..(152331)
; OTHER INFORMATION: n = A,T,C or G
US-09-128-155-16

Query Match 10.2k; Score 140.6; DB 3; Length 152331;
Best Local Similarity 32.8k; Pred. NO. 2.1e-27;
Matches 324; Conservative 166; Mismatches 426; Indels 71; Gaps 6;
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Db 71345 gcaatcctgctag----agagatlagcatgactaaaggagccaaagtgctaatlca 71400
Qy 597 rgayaaygnarayaathsnltgacnltgycaarmgnswmsncarcaruatnha 656
Db 71401 agaaatglttaaaagcccttgtaggcatctcagatctatgaagagagccctccatc 71460
Qy 657 nylngntlmgm-----nccmngmnaartglttyltyggnacngnccngnwnsny 709
Db 71461 acaagtgcaagaggtlgtgctacaaagccagaggttltatggccannccagggccac 71520
Qy 710 tntgtygtnccarccmngayltynglncnngnngnngnnaa----- 755
Db 71521 actgctatgacagccttgtagaacctgctgcccgaatccagccactctgctgctcc 71580
Qy 756 -----ywsngctgtngcnwsnagarngcnwnc 785
Db 71581 acccttgctcaaaagggccaaagatagacttgacacacgctcccgaggccaaagcca 71640
Qy 786 naarcnltgcaarylncnswngnngnngnrcnngntngnngnaaraarwsnmgath-- 843
Db 71641 taagccttggtgttccatgtgttgaagcctgtagcagtgccagaaatgcaagatgag 71700
Qy 844 garytngggarccnccnathmngntlycaraarathaygnaaycngtlyagccmgn 903
Db 71701 ggaagcttggaacttccacctaataatlcagagagatgltcagaacacctagttccca 71760
Qy 904 caraatlygcnngnngnngnswntggnacnswngnngnngnngntnglncaraar 963
Db 71761 cagaagcatgataagagggcagagcccttgcaaggaaacctctataggaatgcaag 71820
Qy 964 gnaayltnngntlggagrcnccncaymngnngnngnngnngnngnngn 1023
Db 71821 gaaatgtggtgtgagctccacacatgltccacactggtgagcactggtgatalct 71880
Qy 1024 gtnmngnwnscnccnswsmngnylncaraarngmngnwnacngywsnytlcar 1083
Db 71881 gtggaatggggctgcgcctccacagaccagaatgtagatgactgtagcagctgcgac 71940
Qy 1084 caytlnccngaraarwnacnngayncartlygarcnngtnaargcnngnngatgar 1143
Db 71941 cctgagccttgaaagctgtagcagcactcaactcaacccaatgataagcaactggct 72000
Qy 1144 wsnlncnctayaaracnngntngnngarylnaacaaracnngntngnathlaytyn 1203
Db 72001 ac--tccagaggagccacagagcgaggtgtgcttagagccttgaggagcctccct 72058
Qy 1204 caytgcaygaayltyngayltnmngcaygngtnaarmngnayaayltyngnngntm 1263
Db 72059 aaccagcttgcaagacatggaatcaa-----agatcatgtgcaagctttaa 72106
Qy 1264 tlygatygcncnngnntlymgnacntlayatggncnnglncnngntlytyngn 1323
Db 72107 ctaaatgttttccctgcacatlcagccttgtyggagccgtgctttttttttt 72166
Qy 1324 tlytycncntlygnaacngnngntlyac 1352
Db 72167 ttttttttggcacaagtglttgac 72195

RESULT 6
US-08-687-080-59
; Sequence 59, Application US/08687080
; Patent No. 5965427
; GENERAL INFORMATION:
; APPLICANT: Gregory Dolganov
; TITLE OF INVENTION: Human RAD50 Gene and Methods of Use Thereof
; NUMBER OF SEQUENCES: 175
; CORRESPONDENCE ADDRESSES:
; ADDRESSEE: Dehlinger & Associates
; STREET: 350 Cambridge Avenue, Suite 250
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
```


Oy	716	gysgtnearccmngmngayltngltnccttygtlncocngtnaaywsngcngltngcncsngar	775
Db	676	gayaibgngcnylntbgcaywsnaartlaygarwsngayacnltngarcagayylngar	735
Oy	776	gngcnwncnnaarccntbgcayltncocwsngcngltngarcngltngngcncnaaraarw	835
Db	736	mgnylnttlycargarylmgngcncnlyntlytnaayccncayactaygtlmgmgngcn	795
Oy	836	snmgnaathgarltntgggarccncnaltmgntlycaraaraalthayagnaaayccntgga	895
Db	796	ylncaymgncaylaygngcncngarytnathgayaltlmgngcncnaltbcncngcncaylt	855
Oy	896	tyccnmncpcraartlytcngcngltngngltngngmsnsgntgmgngacnwsngcngmgng	955
Db	856	ylngngngaraaraaynlyngcncarwsntbggtlnaaathyltngaycngntlytncntly	915
Oy	956	tncaaraaggnaaayltngltngltgggarccncncocncaaymgngltncocnsngngcncnsw	1015
Db	916	yltnaaraaraltbcngcngarygtynacnaaraalthatbgarctncarccayltggarccngar	975
Oy	1016	snmgngcngltmgmgmgwshcncncncnswsnstmgnyltncaraaragmgngnswnacngayw	1075
Db	976	aaryltnaaryltnagaragcngaracntlytlyacntayltngngyltngcngtncncncn	1035
Oy	1076	snyltnarcaaygtlncocngaraaarsnacngayacncarгыarcngltngarocngcng	1135
Db	1036	gncncncncnswntlytggaraaraarylnaaygtlnatbgmgncncncaagaygmngngar	1095
Oy	1136	gnatg 1140	
Db	1096	garly 1100	

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RESULT 13
US-09-351-414-3
: Sequence 3, Application US/09351414
: Patent No. 6265199
: GENERAL INFORMATION:
: APPLICANT: Sheppard, Paul O.
: APPLICANT: Balindur, Nand
: APPLICANT: Delisher, Theresa A.
: APPLICANT: Bishop, Paul D.
: TITLE OF INVENTION: DISINTEGRIN HOMOLOG
: FILE REFERENCE: 98-29
: CURRENT APPLICATION NUMBER: US/09/351,414
: CURRENT FILING DATE: 1999-07-09
: NUMBER OF SEQ ID NOS: 13
: SOFTWARE: FastSeq for Windows Version 3.0
: SEQ ID NO 3
: LENGTH: 2088
: TYPE: DNA
: ORGANISM: Artificial Sequence
: FEATURE:
: OTHER INFORMATION: zdm1l amino acid degenerate sequence
: FEATURE:
: NAME:KEY: variation
: LOCATION: (1)...(2088)
: OTHER INFORMATION: n is any nucleotide
: FEATURE:
: NAME:KEY: misc-feature
: LOCATION: (1)...(2088)
: OTHER INFORMATION: n = A,T,C or G
US-09-351-414-3

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[illegible]

QY	250	aaygcncernngayylnaargarcnatggngntlttgcttgytmgngtncngnnaay	309
Db	791	aytlnatcawsmgmgtlnactltcaytlayaarngwnsmnltwnstlaytlttgargng	850
QY	310	taytawsnwenttgygylngargtlncaycaytgyaaagtlnaayaltbtggyltnatmgm	369
Db	851	tnctywsnmgnacmmgngngtngngtlnaaygartaayagnltncnatgtcngtngcnc	910
QY	370	aarcathtyngcnaayargarathtwsnaatcarcatwsnatcatgargtlnacn	429
Db	911	argtntwtwnscarwsnytngcncatcaaytltngnathcartggargcwnsmngna	970
QY	430	tgggtntlunylnaarngntltysnttyaltlmngargcngarcayaarwsnmgartaaY	489
Db	971	arccnaarttgyaytgyacngarwstbtggngngnttgyalttgargaracngngtlnw	1030
QY	490	ytncaycengayaygnathaararaaraaaycctltysngargsgnaarttyaartln	549
Db	1031	sncaYwsnmgnaatlttwsnaarttgywsnathtyngartatmgngaytltYtncarnng	1090
QY	550	gongcngarathtgyalttgyaaYgaargartynaaycncgcargayaaYgngnar	609
Db	1091	gngngngngcttgytnttlyaaYmgncnacaaraytntltgarccnaengarttgygna	1150
QY	610	aayathwsntggacnttgcarmgnwsnscarcatwsnathtaarwsnyltngcngtmgm	669
Db	1151	aygnttaytngta--rgcngngargarttgyaytgyngnttccaygtngarttgyayg	1208
QY	670	cmmgmngnaarttggtttgyggnacngcngcngnwsnyhtlttgytltcarccmmgn	729
Db	1209	nyltntgytgyaaraarttgywsnyltwnsnaayggngcncaytgywng--aygncnttgyt	1267
QY	730	gaytYngntcncnttgytngcngtlnaaywsngcngtngcngngargcngcwnsncaar	789
Db	1268	gyaayaaYacwnsttgytnttlycatcccmgmngntatgartgymngnaycngtlnaayg	1327
QY	790	cnttgcarytncnwsnngngtngarcngtngngcnaaaraarwsnmgnathtargtln	849
Db	1328	artgyayathacngarttaytgcngngnagaywsngntcarttgcncncaayYlncaya	1387
QY	850	tgggarccncnathmgnttlycaaraathtayggnaayccttsgatcmmgncataraar	909
Db	1388	arcargayngtlaygcttgyaaYcatraaycargymngnttgytlayaayggngarttgyaara	1447
QY	910	ttygngtngngtngnswnsnnttgmgnacnwsngcmmgngtngtncaraargnaay	969
Db	1448	cmmgngaaYacarttgycatatayaltbtgggnacnaarjcngcngngnswngayaarttYt	1507
QY	970	gtngnttggarcncncncaYmgngtlnccnwsnngngcncnwsnmgmngcngtmgm	1029
Db	1508	gytlaygaaartynaayacngargnagncngaraargna----aytgygngaaargayg	1562
QY	1030	mgwnscncnswnswnmgnYtncaraargmgnmgnwsnacngaywsnyltncarcaygln	1089
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QY	1090	cngnraarwnaengayacncarttgcarcncngtlnaaycngcngngatgtgargngtln	1149
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QY	1150	cntlayaaraengtngtngcngartYtnacnaaraacngtngnathtaytYtntlncaYtY	1209
Db	1683	ycaYargtmngngtlnathtgaytgywsngngngncaygtngtntngaytgyaocnga	1742
QY	1210	caygaYtngayttnmgncaygngntnaarmngnaycaytYtgyngcnytmngnttgyay	1269
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OY		471	wsnahcargatgctnactgggttgtytlnuaargcntlywsntlyatactmngaryengar	471
OY	412	412	wsnahcargatgctnactgggttgtytlnuaargcntlywsntlyatactmngaryengar	471
Db	554	554	ACGATTTCAAGAGGTAACTTGGGTGCTGTTAAAAGCATTTAGTTTATAAAGGACAAGG	613
OY	472	472	cyaarwsmwswgsjaraaylncaycnscnaayaagfnahaaaraaaraaycncftysn	531
Db	614	614	CATTGAAGAATTGGAAAAATTTTGCGCCTGCACATGCAATAGAAAGAAATAATCCCATTTCT	673
OY	532	532	gargnnaartlyaaaryltngcngcngaratlyathalyabyaaarylaarylnaaylnaay	591
Db	674	674	GAGGAGAAAATCAACTGGCTGCAGAAAATTTTCATATCTAACACAGAGTAATTAATGTATAT	733
OY	592	592	cnccargayaayygnrgaraayahtwsntngacntyccarmgnwsmwscarcarsnth	651
Db	734	734	CCCCAAGACAAAGGGGAAAAATGCTTCACAGGGGCATCCAGAGGCTTCACAGAGGCCCTC	793
OY	652	652	aarwanlytngcnygmncnmnmng---naarlgtyltvygnaacngncnsgnwsn	708
Db	794	794	CCATCACAGGCTCGCAGGCGCTGAGAGGAAAAAGTATCTCTTGCGGCCAGGCCAAGGATCC	853
OY	709	709	yntlytygtlrcarccmmngayylngtncntlygtfncnnglnaaywsngc-----	761
Db	854	854	CCATCTGTGTGCACCTCGGGACTTGGTGCTTGTGTGCCAGCGTCACACCATGTGCT	913
OY	762	762	-----ngtncwnsrgarqngcngncncaarecntgy	795
Db	914	914	GAAAGGGCCCAACATAGAGCTCGGATTTGTGGCTTCAGAGGGTCGCAAGCTGTAGGCTTGG	973
OY	796	796	carylncnwsmnsngnlngarccngtlngngcnaaaraawsmngnatlhgaryltnbggar	855
Db	974	974	CAGCTTCCATGTGGTGTGAGGCTCCAGTGCACAGAAAGTCAAATAATTTGGGTTGGGAA	1033
OY	856	856	ccnccnatlmgtlycaraaraethaygnaaycctygatgccmmngcaraatlytcn	915
Db	1034	1034	CCTTCGCTTAGATTTTCAGAAAGATGTATGGAAATGCTTGGAATGCTTCGAGCAAGTCTCT	1093
OY	916	916	gtngngntlmgwsmwsntngymnancwnsngnmngtlngtcaraaarygnaayltnng	975
Db	1094	1094	GTAGGGGAGGGGCCCTCATNGAGNAATCTCTGTAGGGCAGTCAACAAGGAAATGTGGGA	1153
OY	976	976	tgggarccnccaymngtlncwnsngngcncwnsmwmngncngtlmngmnwsn	1035
Db	1154	1154	TGCGAGCCCAACACAGAAATCCTACTGGGGCACCGCCTAGTGAGAGCTGTGAGAAGGG	1213
OY	1036	1036	ccnccwnsmwmngnylncaraarygmngmwncnagwywnylncarcayltnccngar	1095
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OY	1096	1096	aarwanacngayacnarctlycarccnglnaargncngngnatlwgrwsnltncntlay	1155
Db	1274	1274	AAAACGGCAGA----CAACACAGGCCCATGAAAGAGCAGTACAGGACAGAGGCTTACCCTGC	1329
OY	1156	1156	aaracngtlnglngarylnaanaaracngtlngnatlhaylynlncayltygayay	1215
Db	1330	1330	AAAGCCACAGGGGGCGGAGTCCCAAGATCAATGAGGAAACCCACTCTTGATCAGATGAC	1389
OY	1216	1216	ytngayglumngcayagnnglnaarnngnayceytltyvgngcngytmngtltyagyecyn	1275
Db	1390	1390	CTGGATGTGAGATTGGAGTTAAAGAGATCATTTTGGAGCTTTAAAGATTTTACGTGCCCC	1449
OY	1276	1276	acngnltlymaachtlayatygnncnft-nccnylnltlytygnncarltytycncnt	1334
Db	1450	1450	TCTAGATTTCAACATCTGCATGGGGCGCTGAGCCCTTGTGTTTACCAATTTCTCCATT	1509
OY	1335	1335	yggnaacngcngntlyacncaartylyntayltncayltgat	1376
Db	1510	1510	TGGAACAGCTGTGTTTACACAAATACCTGTACCCCATTTGTAT	1551

DEFINITION	Homo sapiens, similar to hypothetical protein FLJ14058, clone IMAGE:3831313, mRNA.
ACCESSION	BC004496
VERSION	BC004496.1 GI:14709139
KEYWORDS	HTC.
SOURCE	human.
ORGANISM	Homo sapiens
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
AUTHORS	1 (bases 1 to 2009)
TITLE	Strausberg,R.
JOURNAL	Direct Submission Submitted (12-MAR-2001) National Institutes of Health, Mammalian Gene Collection (MGC), Cancer Genomics Office, National Cancer Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590, USA
REMARK	NIH-MGC Project URL: http://mgc.nci.nih.gov
COMMENT	Contact: MGC help desk Email: cga@bs-research.nhl.gov Tissue Procurement: ATCC/DCTD/DPF CDNA Library Preparation: Rubin Laboratory DNA Sequencing by: The I.M.A.G.E. Consortium (LLNL) http://www.systemsbio.org contact: amadan@systemsbio.org Anup Madan, Rachel Dickhoff, Jessica Fahey, Stephanie Ford, Julia Greene, Mark Kellerman and Anuradha Madan

Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: http://image.llnl.gov/Series/IRAL.Plate: 14 Row: d Column: 7
This clone was selected for full length sequencing because it passed the following selection criteria: Hexamer frequency ORF analysis

This clone has the following problem: frame shifted.
Location/Qualifiers

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1..2009
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/db_xref="taxon:9606"
/clone="IMAGE:3831313"
/lisue_type="Skin, melanotic melanoma."
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/lab_host="DH10B-R"
/note="Vector: pOTB7"

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 Best Local Similarity 45.2%; Pred. No. 1.6e-91;
 Matches 452; Conservative 214; Mismatches 321; Indels 13; Gaps 3;

Qy	390	rargarathwnaarncarcawsmathcarqarjtnacongngnylnynnaarqntt	449
Dd	489	aabagaaatatttttagacacacaaagaacttcmaagtggacttggtttaaggacatt	548
Qy	450	yssnttlialtmngarqcnrgarcayaaasmsnsgnaraaaylncaycngngaayqinat	509
Dd	549	cagatttttaagagaaagcagacataaaaagttcacaaaaatttgccagctgcacaatgat	608
Qy	510	baaraaraaaycncntlywngarqnaartlyaarylngcngcngarathtgyathtg	569
Dd	609	agagaaagaaamaaacatttttttgagcagacaaattccaagccacctgaatttgcatpaa	668
Qy	570	yaaygarqarylnaaygcncaicagayaaygngaraayathwnatgaactglyc	629
Dd	669	tAACAAAGAAAGCGCATGTTAATGCCCAAGACAAGGAAGAAGCACTCCAGACACTATCA	728
Qy	630	rngunswnscncarwsnatnarwsnytnrcvgmngncmmgmnaarttgytlyt	689
Dd	729	GAGGCTTCACAGCAGCCCCCTCCGTACACAGGCCACAGAGGCTTAGGAAAAATGGTTTTCC	788
Qy	690	ygnacnngnccngnwnsynt-----gltcyglncarcemngngaylntglncc--	740

Db	997	TTGTAT	902	EST	11
Db	997	TTGTAT	902	EST	11
RESULT	4				
BI488505					
LOCUS					
DEFINITION					
ACCESSION					
VERSION					
KEYWORDS					
SOURCE					
ORGANISM					
REFERENCE					
AUTHORS					
TITLE					
JOURNAL					
COMMENT					
FEATURES					
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BASE COUNT					
ORIGIN					
Query Match					
Best Local Similarity					
Matches					
493					
2					
553					
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613					
122					
672					
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732					

Db	Accession	Gene	Species	Chromosome	Position	Orientation	Length	Quality	Location
Db	242	CTTGGTGGCCCTGTGCTCCAGTCCAGCTGTACATGAAAGGGGCCAACGTAGAGCTCG	Human	1	1050	+	1050	High quality	1050
QY	759	ngcngtngcwsnagarargngcwsnccnaarccnhtgcaaytlncnwsngngtngarc	Human	1	1050	+	1050	High quality	1050
Db	302	GGCTGTGGCTTCAGAGGGTGGAGCTTCMACTTTGGAGCTTCACAGTGTGTGGAGCC	Human	1	1050	+	1050	High quality	1050
QY	819	ngtngngcnaaraarwsmngnathgargtltggagrcnccnahtmgntlycaaraat	Human	1	1050	+	1050	High quality	1050
Db	362	TACAGGTCCACAGAAGTCAAGAACTGAGGTTTGGAGACCTTGCTGATTTTCAGAAAT	Human	1	1050	+	1050	High quality	1050
QY	879	htaygnaaaycctngatlycncmngncaraatlycngtngngtngngnwsnawtngm	Human	1	1050	+	1050	High quality	1050
Db	422	GTTTGGAATAGCTGGATGCCAGCCAGCAAAATTTTGCTCAGAGAGTGGGCCCTTATGAG	Human	1	1050	+	1050	High quality	1050
QY	939	nacnwsngcmngtngtncaraargnaaytngngtnggagrcnccnaymngtnc	Human	1	1050	+	1050	High quality	1050
Db	482	AACCTCTGAGACATGCGAAGGAAATGTGGATGGAGGCCCCACAGAGAGTCCA	Human	1	1050	+	1050	High quality	1050
QY	999	nwsngngcncnwsnwsnmngngtngmngmwsnccnccnwsnwsnmngnlyncara	Human	1	1050	+	1050	High quality	1050
Db	542	TACTGTGGGACACTGCTTACTGTGGAATGTGAGAGACAGCCAGCTCTCAACACCCAGAA	Human	1	1050	+	1050	High quality	1050
QY	1059	rgmngnwsnawcngaynylnarcaaaytncnngaraarwsnawcngayncartgca	Human	1	1050	+	1050	High quality	1050
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QY	1119	recngtlnaargcngcngnagatgargtngtncntayaraacngtngtngcngartnac	Human	1	1050	+	1050	High quality	1050
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QY	1179	naarcngtngnathgaytngtncaytlygcaaytngagfymngncayngntaa	Human	1	1050	+	1050	High quality	1050
Db	722	CAGAGCAGATGGGAGACCACTCTTGATCAAGCTTGACCTTGATGGAGAACTGAGTAA	Human	1	1050	+	1050	High quality	1050
QY	1239	rngnagaycaltlygngcnytl-nmgntlygalytygcacnngntlymnahtayalg	Human	1	1050	+	1050	High quality	1050
Db	782	AAGAGATGCTTTTGGAGCTTAAACATGATGACCTGCGCTGATTCAGACTTCATVG	Human	1	1050	+	1050	High quality	1050
QY	1298	gncngt 1304	Human	1	1050	+	1050	High quality	1050
Db	842	GGGCGCT 848	Human	1	1050	+	1050	High quality	1050
RESULT	5	BM472108	1050 bp	mRNA	linear	EST 05-FEB-2002			
LOCUS		BM472108							
DEFINITION		AGENCOURT_6465359 NIH_MGC_72 Homo sapiens cDNA clone IMAGE:5539381							
ACCESSION		BM472108							
VERSION		BM472108.1							
KEYWORDS		EST.							
SOURCE		human.							
ORGANISM		Homo sapiens							
REFERENCE		Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;							
AUTHORS		Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.							
TITLE		NIH-MGC http://mhc.nci.nih.gov/.							
JOURNAL		National Institutes of Health, Mammalian Gene Collection (MGC)							
COMMENT		Unpublished (1999)							
		Contact: Robert Strusberg, Ph.D.							
		Email: cga@rs-remail.nih.gov							
		Tissue Procurement: ATCC/DCT/DTF							
		cDNA Library Preparation: Life Technologies, Inc.							
		cDNA library Arrayed by: The I.M.A.G.E. Consortium (LLNL)							
		DNA Sequencing by: Agencourt Bioscience Corporation							
		Clone distribution: MGC clone distribution information can be							
		found through the I.M.A.G.E. Consortium/LLNL at:							
		http://image.llnl.gov							
		Plate: LLAM12233 row: m column: 14							
		High quality sequence stop: 658.							
		Location/Qualifiers							
FEATURES		1..1050							
source		/organism="Homo sapiens"							

Db	Accession	Version	Keywords	Source	Organism
Db	420	CATGAGAAACCTCTGCTTAGGGACAGTGTGGAAAGGAAATGTGGGGTTTGGAGCCCGCACACA	479		
Oy	992	gnftrcnwsmngngcncncnswmsnmngcngtmmgmgnwscncncnswmsnmngny	1051		
Db	480	GTGGCCCTGCTGGGGCCATGCTCCTAGTGTGAGAGGCTGTGTAGAAAGGGCCACCATCTCTCCAGAC	539		
Oy	1052	ttcacaatgaatgmgmsnscnaysnynlnarceaytcttcnagarataarwnaenayaenc	1111		
Db	540	CCGAGAAATGTAAGATCCATCAACACTTGGCACTGTGGCACTTGGAAACCCGACAGACATC	599		
Oy	1112	artgycaarcnngtlnaarycngcngnaltvgarwsmngtncnltayaaraengtngtngcng	1171		
Db	600	GACACACCCCATGATAAAGACAGCTGGAGAGGGAGGCTATACCTCGCAAAACACAGGGCGAG	659		
Oy	1172	arythacnaaatacngtngnathaytynhncaytgycaaytngaytngaytmgncayg	1231		
Db	660	AGCTGCTCTTAAGACCATGGGGAACCCACTCTTGGACACACATGACCTGGATATGAGACTTG	719		
Oy	1232	gngtlaaamgmngaycaaytctgyngcngnytmngtlygaytgycaacnagntlymnaent	1291		
Db	720	GAGTCAAAAGGAGATCATTTTGGAGCTTTGAATTTGATGATTCGCCCGCTGGATTTTCAGACTT	779		
Oy	1292	ayaatgngcngt-nccnytnbytlytngncarttytctytcntlytngnagcngtntly	1350		
Db	780	GTGTANGGCTGATAACCCCTTTGTTTGGGCAATTTCCATTTTGATGCTTCAATTT	839		
Oy	1351	acncartgtytnhtaytncaytgyat	1376		
Db	840	TACCAATACCTTACCCCTTAGGGAT	865		
RESULT	7				
LOCUS	BM457166	898 bp	mRNA	linear	EST 05-FEB-2002
DEFINITION	AGENCOURT_6411690 NIH_MGC_92 Homo sapiens CDNA clone IMAGE:5583427				
ACCESSION	BM457166				
VERSION	BM457166.1	GI:18506206			
KEYWORDS	EST.				
SOURCE	human.				
ORGANISM	Homo sapiens				
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;				
AUTHORS	Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.				
TITLE	1 (bases 1 to 898)				
JOURNAL	NIH-MGC http://nigc.nci.nih.gov/ .				
COMMENT	National Institutes of Health, Mammalian Gene Collection (MGC) Unpublished (1999) Contact: Robert Strausberg, Ph.D. Email: cga@bbs-renal.nih.gov Tissue Procurement: ATCC cDNA Library Preparation: Life Technologies, Inc. cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL) DNA Sequencing by: Agencourt Bioscience Corporation Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: http://image.llnl.gov Plate: LLNL12346 Row: h Column: 20 High quality sequence start: 9 High quality sequence stop: 713. Location/Qualifiers 1..898 /organism="Homo sapiens" /db_xref="taxon:9606" /clone_image="5583427" /clone_id="NIH_MGC_92" /tisue_type="embryonal carcinoma, cell line" /lab_host="DH10B (phage-resistant)" /note="Organ: testis; Vector: pCMV-SPORE6; Site:1: NotI; Site:2: SalI; Cloned unidirectionally; oligo-dT primed. Average insert size 2.5 kb. Library enriched for full-length clones and constructed by Life Technologies. Note: this is a NIH-MGC Library."				
FEATURES					
SOURCE					

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OY	622	acnlyearmgnwsmwsmncarcarwsnaetharwsnylngntlgmngcmgmng---n	678		
Db	95	TCATGTCCAGAGACCTTCATATGCGACGCCCTCCCTCCATCACAGGCCACAGGCCACAGAGGAAA	154		
OY	679	aartgltlytgygnaacngnccngnwsnytnlytgytlnacarccmgnayxytngtn	738		
Db	155	AAGTGGTTTGTGGGCTGGGCCACAGGTCCTGTCTGTCTGTGTGACGACTAGGACCTTGSTG	214		
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OY	766	gcnwsngargngnccnwsncnnaarcntggcarytlnccnwsngngtltgarcnngtn	825		
Db	275	GCTTAGAGGGGTAGAGGGCCCAAGCATTTGGCAGCTTCATGTGATGTGAGTTCCTGTGGGT	334		
OY	826	gcnnaararwsnmgathgargtlnbggargcncnccnatlmngtlycaratrbhargn	885		
Db	335	GCAGAGATGTCAAGATTTAGATTTTGGGAACTCTGCTTACATTTTCAACAATATATATGGA	394		
OY	886	aaycnnlytgatccmngncararctlycngtngngtngngtnwsmnltgmgmncnwsn	945		
Db	395	AACACTGGATGTCACAGGCAAAAGTTTCTGTGAGGGGACAGGCCCTCATGGGAAACCTCT	454		
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OY	1006	gcnccnwsnwsnmgngcngtmgmgngnccnccnws-----n	1044		
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Db	574	GGTAGAACCCCAAAATGTGTAGATCAACACCAACTTGTGCTTGTGCTGGGAAAAACTTACA	633		
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Db	634	GACACTCAACATCAACCTCTGAGAAAGACAGCAGAGGAGGCTGTACTCTGCAAAACCCACA	693		
OY	1165	gtnccngarytlnaacaaraengtngnathelytlnylnccaytlycaygaytlyngaytn	1224		
Db	694	GGGGGGGAGACTGCACAAAGCCATGGGAACCTATCTCTTCACATCACATGACCTGTGATGTG	753		
OY	1225	mgncaygngtlnaaraargaycarytlygngcngnylmngtlygargtlycncnccngntly	1284		
Db	754	AGACATGGGAGTCAAAAGAGACATCATTTATGTGAGCTTTAAATTTTGACTGTGCCACCTGATTT	813		
OY	1285	mgnaonlayatlgngcngnt--nccnytnlytlytgnacartlytlycnnltlygnaacngc	1343		
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DEFINITION	Pan troglodytes DNA, clone:PTB-085H08.F, genomic survey sequence.				

Accession	KeyWords	Source	Organism
AC086951	AC086951.1 GI:16638753	GSS; GSS (genome survey sequence).	
		Pan troglodytes male lymphoblast DNA, clone_11b:PTB Chimpanzee Male	
		BAC library clone:PTB-085H08.F.	
		Pan troglodytes	
		Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;	
		Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Pan.	
REFERENCE	1 (sites)		
AUTHORS	Fujiyama,A., Hattori,M., Toyoda,A., Taylor,T.D., Yada,T.,		
TITLE	Tokoi,Y., Watanabe,H. and Sakaki,Y.		
REFERENCE	BAC end sequences of library PTB		
AUTHORS	Unpublished		
TITLE	2 (bases 1 to 717)		
JOURNAL	Fujiyama,A., Hattori,M., Toyoda,A., Taylor,T.D., Yada,T.,		
	Tokoi,Y., Watanabe,H. and Sakaki,Y.		
	Direct Submission		
	Submitted (02-AUG-2001) Aaso Fujiyama, The Institute of Physical		
	and Chemical Research (RIKEN), Genomic Sciences Center (GSC);		
	1-7-22 Suehiro-cho,Tsukumi-ku, Yokohama, Kanagawa 220-0045, Japan		
	(E-mail:chimpbes@sc.riken.go.jp, URL:http://hsp.gsc.riken.go.jp/,		
	Tel:81-45-503-9111, Fax:81-45-503-9170)		
	Clones are derived from the chimpanzee BAC library PTB This BAC end		
	was generated during the Red process and may have higher chance of		
	clone tracking errors.		
COMMENT	PRIMERS		
	Sequencing: -21M13		
FEATURES	LIBRARY		
source	Vector : pKS145		
	R.Site 1 : SacI		
	R.Site 2 : SacI.		
	Location/Qualifiers		
	1..717		
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	/db_xref="taxon:9598"		
	/clone="PTB-085H08.F"		
	/sex="male"		
	/cell_type="lymphoblast"		
	/clone_11b="PTB Chimpanzee Male BAC library"		
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Query Match	22.7%	Score 313,	DB 12,
Local Similarity	49.1%	Pred. NO. 1.7e-60;	Length 717;
Matches 299;	Conservative 113;	Mismatches 195;	Indels 2;
		Gaps 1;	
744 ygttcngttnaaywngngntgcnwngarzgngcnwscncaarcentgtgacrttcc	803		
89 cctccggcgagctcagcgtgtgcttcgagggctccaaagccttaagccttgccacgttcc	148		
804 nwsngngtngarncngtngngcnaaraarwsnmgnaathgargtntggarcnccnat	863		
149 acatggtgttaagccttgatgcagtcacagaaatcaagaaactggcggttggaacctctgcc	208		
864 hmgnttcaaraarathayggnaaaycmttgatgctcmngncaraarttygcngtngntg	923		
209 tgattttcagagatataatgttaataccttgatgccacagaaattttgctgtaggga	268		
924 nggnwsnwtngmgnaacnwsngcmgngtngtncaraargnaaygtngnttgarc	983		
269 gggggccttcagtaaaaacacctgtcgaaggcactgtggagagaaatgtgggtcagaccc	328		
984 nccncaymngtncnwsngngnccmwsnsmngndcngtngnmgnwsnccnws	1043		
329 atccacacagagctcctactactggggcaccgcccttagtgagacctgtgaaaagtggccctatt	388		
1044 nwsnmngtncaraargnmgnaacnagaaywnylncarcaygttncngnaraarwsnac	1103		
389 ctccagaccctcagaaatgtatgtccatgcacagcttgacacctgcagaaagccac	448		
1104 ngaycncatrttyccnctngttnaaygcngcngnatgtgarnsngtnccttayaaracngt	1163		

Db	449	AGACATCAATGCCAGCGCTGTGAAGACGACTGGAGGGAGACTGACATGGAAGACAC	508
Oy	1164	ngtgcngarytfnacnaaracngtngnathlaytlnylncaeytgcaygalytngayt	1223
Db	509	AGGGGACGAGCTGCCCAAGACCATGGGAACCCACCTCTTGATCATCATGACCTGGAGT	568
Oy	1224	nmgeaygngtfnarmngaycaytlygngngcnytmngntlygalytgcacnagntt	1283
Db	569	GAGACATGAGCCAAAGGAGATCATTTGGAGACTTTAAGATTGACTTCCTGCTGGANT	628
Oy	1284	ymgancntaycagggcngct--nccnlytlnlytngarycartytlycnclytgnacn	1341
Db	629	TGACAGCTTGCAATGGGCGCTGTGAGCCCTTTTTTTGGCAATTTCCGCCAATTGGAGT	688
Oy	1342	gcngntty 1350	
Db	689	GCGTTAATT 697	
RESULT	9		
LOCUS	B1754555/c	736 bp	mRNA linear EST 25-SEP-2001
DEFINITION	603023384P1 NIH_MGC_114 Homo sapiens cDNA clone IMAGE:5194028 5',		
ACCESSION	B1754555		
VERSION	B1754555.1	GI:15746133	
KEYWORDS	EST.		
SOURCE	human.		
ORGANISM	Homo sapiens		
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;		
AUTHORS	Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.		
TITLE	1 (bases 1 to 736)		
JOURNAL	NIH-MGC http://mgc.nci.nih.gov/ .		
COMMENT	National Institutes of Health, Mammalian Gene Collection (MGC)		
	unpublished (1999)		
	Contact: Robert Strausberg, Ph.D.		
	Email: cgabs-remail.nih.gov		
	Tissue Procurement: Life Technologies, Inc.		
	cDNA Library Preparation: Life Technologies, Inc.		
	cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)		
	DNA Sequencing by: Incyte Genomics, Inc.		
	Clone distribution: MGC clone distribution information can be		
	found through the I.M.A.G.E. Consortium/LNL at:		
	http://image.llnl.gov		
	Plate: LLM11485 Row: 9 Column: 21		
FEATURES	High quality sequence stop: 733.		
source	Location/Qualifiers		
	1..736		
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	/db_xref="taxon:9606"		
	/clone="IMAGE:5194028"		
	/clone_lib="NIH_MGC_114"		
	/lab_host="DH10B"		
	/note="Organ: brain; Vector: pCMV-SPORT6; Site.1: NotI;		
	Site.2: EcoRV (destroyed); RNA source anonymous pool of 6		
	male brains, age range 23-27 yo. Library is oligo-dT		
	primed and directionally cloned (EcoRV site is destroyed		
	upon cloning). Average insert size 1.5 kb, insert size		
	range 1-3 kb. Library is normalized and enriched for		
	full-length clones and was constructed by C. Gruber		
	(Invitrogen). Research Genetics tracking code 019. Note:		
	this is a NIH_MGC Library."		
BASE COUNT	163 a 196 g 185 g 192 t		
ORIGIN			
Query Match	22.3%, Score 306.6; DB 10; Length 736;		
Best Local Similarity	44.1%; Pred. No. 5.1e-59;		
Matches 306; Conservative 122; Mismatches 266; Indels 0; Gaps 0;			
Oy	659	tngcttgmgncemgmngnaarigtgttytlygngacngcngcngmsnytntgtytg	718
Db	694	tggcttgagcccaagctcccatgtgttgtagcagcttgagacttgcctgcctgctccca	635

ACCESSION	AG064424
VERSION	AG064424.1 GI:16616226
KEYWORDS	GSS; GSS (genome survey sequence).
SOURCE	Pan troglodytes male lymphoblast DNA, clone_1lb:PTB Chimpanzee Male BAC library clone:PTB-053H24.R.
ORGANISM	Pan troglodytes
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Pan.
AUTHORS	1 (sites)
TITLE	Fujiyama,A., Hattori,M., Toyoda,A., Taylor,T.D., Yada,T., Totoki,Y., Watanabe,H. and Sakaki,Y.
JOURNAL	BAC end sequences of library PTB
REFERENCE	Unpublished
AUTHORS	2 (bases 1 to 676)
TITLE	Fujiyama,A., Hattori,M., Toyoda,A., Taylor,T.D., Yada,T., Totoki,Y., Watanabe,H. and Sakaki,Y.
JOURNAL	Direct Submission
COMMENT	Submitted (02-AUG-2001) Asao Fujiyama, The Institute of Physical and Chemical Research (RIKEN), Genomic Sciences Center (GSC), 1-7-22 Sueniro-chou,Tsurumi-Ku, Yokohama, Kanagawa 230-0045, Japan (E-mail:climpb@esgsc.riken.go.jp, URL:http://hnp.gsc.riken.go.jp/, Tel.:81-45-503-9111, Fax:81-45-503-9170) Clones are derived from the chimpanzee BAC library PNB This BAC end was generated during the Rsd process and may have higher chance of clone tracking errors. PRIMERS Sequencing: MJ3rev LIBRARY Vector : pKS145 R.Site 1 : SacI R.Site 2 : SacI. Location/Dualifiers FEATURES source 1..676 /organism="Pan troglodytes" /db_xref="taxon:9598" /clone="PTB-053H24.R" /sex="male" /cell_type="lymphoblast" /clone_lib="PTB Chimpanzee Male BAC Library"
BASE COUNT	164 a 165 c 191 g 156 t
ORIGIN	
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Best Local Similarity	47.9%; Pred. No.1.3e-56;
Matches 294; Conservative 117; Mismatches 201; Indels 2; Gaps 2;	
Oy	760 gcngtngcngsnrgatrgngcnmsncncnaarctntggacrytncnswngngtgarcgn 819
Dd	63 GCCATGGCTTCTAAAGGTGCACAGCCTTCAGCGCTTGCGACTCCACATGATTGAAGTG 122
Oy	820 gtngngcnaaarawsmgna lbhargltngbgarcncnccnathmgnttcaraarath 879
Dd	123 GCMAATGCACAGCAAGTCAAGAAGCTGGGGTTTAGAAGCTTAACCTTAAGATTTCAGAGATG 182
Oy	880 taygnaayccnttgcgtgccmmgncaraarttygcngtngngtngngtngwsmwtcgmn 939
Dd	183 TATGGAATGCTGGATGATGCCAGGAGAAATTTCGCCGAGGAGCGGGGCTCTTGAGGA 242
Oy	940 acnwsgncgmngngtngtncaraarqnaaygtngngtggarcncncnaymngtncncn 999
Dd	243 ACCCTTGCTAGGGCATGTGMAAGGAAATGTGGGGTTGGAGTCCACCACCGACGATCCCT 302
Oy	1000 wangngcncnmwasmngngcngttnmgngwnwcncncnmwsmngmtncaraar 1059
Dd	303 ACTGGGGCACCATCTCAAGTGGAGCTGTGAGAAC-AGGCACTGACCTCCAGACCCAGAAAT 361
Oy	1060 ggmgmwsmnacngaywmytlncaarqytlnccngaraarawsmnacngayacncaatgycar 1119
Dd	362 GGATAGTCACTGACAGAGCTTACACCATGTGGCTGAAAAAGCTTAACACCTCAACACGAG 421
Oy	1120 ccngtngaargcngcngnatvgarwsngtncntayaaacnctngtngtngcngarytncacn 1179

Db	422	CCCATGAAAGCATCTGGGAGGGAGGTGTCTACCTGTGCANATCCACAGGGGGAGAGTGGCC	481
Qy	1180	aaracngtngnacthtaytctnlyncaytgycaayaaytngaygtmngcaygngtinaar	1239
Db	482	AAGACATGAGGAGCCACCTCTTGATCAGTGTGACCTGGATGTGACATGAGATCMAA	541
Qy	1240	mngnagcaycttggngcngtctmngtnttgyatgygcncnagngnttymnactataytgg	1299
Db	542	GGAGATCATTTTGTGATGTTTAAAGATTTGACGTGCTCGCTGATTTTGGACATGAGCATG	601
Qy	1300	cgngt-nccncttngtlytngnacttlytccnttlytngnagcngngnttlyacnctg	1358
Db	602	CCTGTAGCCCTCTTGTCTTTGGCCATTTCTCCCATTTTGGAATGGCTGTATTTACCAATG	661
Qy	1359	yytntayttncayt	1372
Db	662	CCTGTACCCCATTT	675
RESULT	12		
AG062447			
LOCUS	AG062447	691 bp	DNA
DEFINITION	Pan troglodytes DNA, clone: PTB-050M20.F, genomic survey sequence.		
ACCESSION	AG062447		
VERSION	AG062447.1	GI:16614249	
KEYWORDS	GSS; GSS (genome survey sequence).		
SOURCE	Pan troglodytes male lymphoblast DNA, clone_11b:PTB Chimpanzee Male BAC Library clone:PTB-050M20.F.		
ORGANISM	Pan troglodytes		
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Pan.		
AUTHORS	1 (sites)		
TITLE	Fujiyama, A., Hattori, M., Toyoda, A., Taylor, T. D., Yada, T., Tokoki, Y., Watanabe, H. and Sakaki, Y.		
REFERENCE	BAC end sequences of Library PTB		
AUTHORS	Unpublished		
TITLE	2 (bases 1 to 691)		
JOURNAL	Fujiyama, A., Hattori, M., Toyoda, A., Taylor, T. D., Yada, T., Tokoki, Y., Watanabe, H. and Sakaki, Y.		
COMMENT	Submitted (02-AUG-2001) Asao Fujiyama, The Institute of Physical and Chemical Research (RIKEN), Genomic Sciences Center (GSC); 1-7-22 Shikui-chou, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan (E-mail:shimbes@sc.riken.go.jp, URL:http://hgp.gsc.riken.go.jp/, Tel:81-45-503-9111, Fax:81-45-503-9170)		
COMMENT	Clones are derived from the chimpanzee BAC library PTB This BAC end was generated during the R&D process and may have higher chance of clone tracking errors.		
PRIMERS			
Sequencing:	-21M13		
LIBRARY			
Vector	: pKS145		
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R.Site 2	: SacI		
Location/Qualifiers			
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/clone="PTB-050M20.F"			
/sex="male"			
/cell_type="lymphoblast"			
/clone_11b="PTB Chimpanzee Male BAC Library"			
BASE COUNT	160 a 167 c 201 g 162 t		1 others
ORIGIN			
Query Match	20.9%; Score 287.4; DB 12; Length 691;		
Best Local Similarity	49.0%; Pred. No. 1.2e-54;		
Matches 301; Conservative 110; Mismatches 196; Indels 7; Gaps 3;			
Qy	744	ytgctngcgttgaatgngcngtngcngtngatgagngcngcncncaarctgtgcarytnc	803
Db	82	CGTCGGCGAGCTCAGGCTGTGGCTTCAGAGAGGTCAAGACCCTTGTGGAGGCTTCC	141

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QY 804 nwsngngtngarcngtngngngcnaaraarwsmngnathgargtntggarcnccnat 863
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Db 1A2 ATGTGCTGTGAGCTGTGAGTGACAGAGAGA----ATCGGGGTTTGGGAACCTCCACT 196
QY 864 hmngtlycaraarathcygnaayccttgatgctcmngncaraartlygcngtngngt 923
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Db 197 TAGATATGAGAAGATGTATGAAAATACCTGGATGCCATGCAAAAAGTTTGGCTGTAGGGTC 256
QY 924 ngngnswntgmgnaacwngcngmgngtngtncaraargnaaygtngtngtggarcc 983
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Db 257 AGGGCCCTCATGAGAACTTCTGCTAGGGCAGCTGTGAGAGGAATGTAGGTCAGAGGC 316
QY 984 nccncaymngtncnswngngngcncswngsmngngngtngmngmngwsmcncnccms 1043
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Db 317 CCCACACAGAGTCCCTACTGGGGCACACACAGTGAGAGCTGTGAGAGAAGGGCCACTGTC 376
QY 1044 nwsnmngtncaraargnmngsmnagaywsnytncaaraytncngngaraarwsmnac 1103
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Db 377 CTCCAGACTCCAGATGGGTAGATCCATCCATGCTTGATCATGTGCTCGGAAAAGCCAC 436
QY 1104 ngayancartlycarcngtngarcngcngngngatggarwngtncntayaraarngt 1163
    |||::|||::|||::|||::|||::|||::|||::|||::|||::|||::|||::|||
Db 437 AGACACTCAATGCCAGCCCATCAAAAGCAGCCGGAAGAGGCTGTACCTACAGAGCCAC 496
QY 1164 ngtngcngaryt-nacnaaracngtngngnathlaytngtncaytlycayagaytngay 1222
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Db 497 AGGGCAGAGCTGTCCAAAGACCATGGAGACCACCCCTTGCTCAGTGCACGTGTTG 556
QY 1223 tnmngcaygngtngnaarngngaycaytlygngcngtngmtngtlycaygycnaengnt 1282
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Db 557 TGAGATATGAGATCAGAGAGATCATGTAGAGCTTTAAGATTTGACCTGTGGTGAT 616
QY 1283 tymnagactlayatggngcngnt-nccnytnlytlytngncartlytlycngtlygnaon 1341
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Db 617 TTGGGACTTGCATGGGGCTGTAGCTCTTCTTTTGGCCATTTCTCCATTGTGAATG 676
QY 1342 gcnngtntlyacna 1355
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Db 677 GCTGTATTTACCCA 690

RESULT 13
AG116938
LOCUS AG116938 720 bp DNA linear GSS 03-NOV-2001
DEFINITION Pan troglodytes DNA, clone: PTB-124K01.R, genomic survey sequence.
ACCESSION AG116938
VERSION AG116938.1 GI:16737457
KEYWORDS GSS: GSS (genome survey sequence).
SOURCE Pan troglodytes male lymphoblast DNA, clone: PTB Chimpanzee Male
        BAC Library clone: PTB-124K01.R.
ORGANISM Pan troglodytes
        Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
        Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Pan.
REFERENCE 1 (sites)
AUTHORS Fujiyama,A., Hattori,M., Toyoda,A., Taylor,T.D., Yada,T.,
        Tokoki,Y., Watanabe,H. and Sakaki,Y.
TITLE BAC end sequences of Library PTB
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 720)
AUTHORS Fujiyama,A., Hattori,M., Toyoda,A., Taylor,T.D., Yada,T.,
        Tokoki,Y., Watanabe,H. and Sakaki,Y.
TITLE Direct Submission
JOURNAL Submitted (02-AUG-2001) Asao Fujiyama, The Institute of Physical
        and Chemical Research (RIKEN), Genomic Sciences Center (GSC), Japan
        1-7-22 Suehiro-cho,Tsukumi-ku, Yokohama, Kanagawa 230-0045, Japan
        (E-mail:chiimpes@gsc.riken.go.jp, URL:http://ngp.gsc.riken.go.jp/,
        Tel:81-45-503-9111, Fax:81-45-503-9170)
COMMENT Clones are derived from the chimpanzee BAC library PTB This BAC end
        was generated during the Rad process and may have higher chance of
        clone tracking errors.
PRIMERS
Sequencing: M13Rev
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LIBRARY
Vector : pKS145
R.site 1 : SacI
R.site 2 : SacI.
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Source
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/clone="PTB-124K01.R"
/sex="male"
/cell_type="lymphoblast"
/clone_lib="PTB Chimpanzee Male BAC Library"
BASE COUNT 181 a 186 c 187 g 165 t 1 others
ORIGIN

Query Match 20.8%; Score 286.2; DB 12; Length 720;
Best Local Similarity 48.5%; Pred. No. 2.3e-54;
Matches 298; Conservative 112; Mismatches 200; Indels 4; Gaps 3;

QY 760 gngtngcwnsngarargngcwnscncaarcntlygcarlytncnswngngtngarcn 819
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Db 110 GCCATGCGCTTCAGATGCTGCACAGCACCAAGTCTTGCTGCACATGACCTTGAAGCT 169
QY 820 gtnngcnaaraarwsmngnathgargtngtnggarcncncaatngtlycaraarath 879
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Db 170 GCGGCTGCACAGAAATGAGATGAGGTTTGAAGACCTCCAGTAATTCAGAAATG 229
QY 880 taygnaayccttgatgctcmngncaraartlygcngtngngtngtngmngsmngngn 939
    |||::|||::|||::|||::|||::|||::|||::|||::|||::|||::|||::|||
Db 230 TATGGAATATGCTTGATGCCAGGCAAGATTTGCTGAGGGAGTGGCTCTCATGAGAGA 289
QY 940 acnwsngcngngtngtncaraargnaaytngtngtgargcncncncaymngtncn 999
    |||::|||::|||::|||::|||::|||::|||::|||::|||::|||::|||::|||
Db 290 ACCTCTGCTAGGCGCAGTGCAGAGGAATATGGGGTGGAGACCCACACAGGGTCCCT 349
QY 1000 wsnngcncnswngsmngngcngtngmngmngwsmcncnccnswsmngntncaraar 1059
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Db 350 ACTGGGG--ATCACCTTAGTGAGAGCTGTGAGAAAGACCCACATCTCTCCAGACCCAGAA 408
QY 1060 ggmngnswnacngaywsnytncaaraytncngngaraarwsmnagayancartlycar 1119
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Db 409 GCTTAATCCACTGACACTTGC--ACCTGTGCTGGAAGAGCTGCAGATCTCAATCAATG 466
QY 1120 cngtncnaarngcngngatlgarwngtncntayaraarngtngtngcngarytlna 1179
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Db 467 CCCATGAAAGCAGCTGGGAGGAGGCTGTACCTGCAGAAAGCAGAGAGGTGAGCTTCC 526
QY 1180 aaracngtngnathlaytngtncaytlycayagaytngaytngmngcaygngtlna 1239
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Db 527 GAGACCATGGGAACCTACTCTTGATGAGGTGACCTGGATGTGAGCATGATTCATAA 586
QY 1240 mngngaycaytlygngcngtngmtngtlycaytlygcnaengntlymgnactaytgg 1299
    ::|||::|||::|||::|||::|||::|||::|||::|||::|||::|||::|||::|||
Db 587 AGAGATCATATTGGAGCTTTAAGATTGACAGCCACACAGCTGGATTTGGACTTGATGGC 646
QY 1300 cngt-nccnytnlytlytngncartlytlycngtlygnaengngtngtlycncart 1358
    |||::|||::|||::|||::|||::|||::|||::|||::|||::|||::|||::|||
Db 647 CCTGTAGCCCTTTGTTTGGCCATTTCTCCATTTCGTAAGTTCATTAACCTAATG 706
QY 1359 yytntaytlncaayt 1372
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Db 707 TCTGTATTCCTCAT 720

RESULT 14
AQ037711
LOCUS AQ037711 623 bp DNA linear GSS 11-JUL-1998
DEFINITION CIR-HSP-2337G10..TR CIR-HSP Homo sapiens genomic clone 2337G10, DNA
        sequence.
ACCESSION AQ037711
VERSION AQ037711.1 GI:3303543
KEYWORDS GSS.
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SOURCE	human.
ORGANISM	Homo sapiens
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo. 1 (bases 1 to 623)
AUTHORS	Adams,M.D., Rounsley,S.D., Zhao,S., Field,C.E., Bass,S., Linher,K., Golden,K., Berry,K., Granger,D., Suh,E., Wible,C., Shizuya,H., Simon,M. and Venter,J.C.
TITLE	Use of a random BAC End Sequence Database for Sequence-Ready Map Building (1998)
JOURNAL	unpublished (1998)
COMMENT	Other_GSSs: CIT-HSP-2337G10.TF Contact: Mark Adams Department of Eukaryotic Genomics The Institute for Genomic Research 9712 Medical Center Dr., Rockville, MD 20850, USA Tel: 301 838 0200 Fax: 301 838 0208 Email: madams@tigr.org Clones are available from Research Genetics (info@resgen.com) . BAC end search page: http://www.tigr.org/tdb/humgen/bac_end_search/bac_end_search.html . Seq primer: M13 Reverse Class: BAC ends.
FEATURES	Location/Qualifiers
SOURCE	1..623 /organism="Homo sapiens" /db_xref="taxon:9606" /clone="2337G10" /clone_1lb="CIT-HSP" /sex="Male" /cell_type="Sperm" /note="Vector: pBeloBAC11; Site_1: HindIII; Site_2: HindIII"
BASE COUNT	160 a 146 c 172 g 145 t
ORIGIN	
Query Match	20.6%; Score 283.8; DB 12; Length 623;
Best Local Similarity	48.1%; Pred. No. 7.3e-54;
Matches	286; Conservative 111; Mismatches 196; Indels 2; Gaps 2
OY	784 cnaarccttgccatgttcncwngngtngarcngtngngcngtnaaaraarwsnmgnath 843
Db	26 CTCAGCTTTGGCAGACTTCACATGCTGTGAGTCTCGCGGATCACCAAGAAGCAGAAACT 85
OY	844 gaigtntg-ggaarcncnathmgnltcacaaraatltagysnaaycentgatgcmmg 902
Db	86 GAGGTGTGAAGAACCTCCACCTACTTTTCAGAAAGTAGTAGGAATAATGCCGTGATCTCCAG 145
OY	903 ncaaraattlycngtngngtnggwswanlytgmgagncwngcnmgngtngtncaraa 962
Db	146 GCAAAAGCTTCTCTCAGGGGCAGGGCCCCAGATGGAGAACCTTGCTAGGGCAGTGGGAA 205
OY	963 rfgnaeytngtngtgygarccnccncayymgntlncwswngngcncswnswnsmngnc 1022
Db	206 GGGAATGTGGGGTTGGAGCCGTGCACAAGAGTCTCTACTGGGCACGTGCTAATGGAGAC 265
OY	1023 ngttmgmgmwsnccnccnswnsnmgnlytncaraarygmngnwsnacngaywsnytnca 1082
Db	266 TGTGAGAGAGGAGGACTTGTCTCTCCAGACCCTTAAGATGATGATGCTGCCGACACTTGCA 325
OY	1083 rcaygttncngaraarwsnacngayaacncartgyarccngtinaargcngtngnatgga 1142
Db	326 TTATTGCACACTGGAAAAAGCCACAGACACTCAACACCAAGCCCATGAAAACAGCCAGGGGG 385
OY	1143 twsngtncnclayaaraengtngtngcngaryttnaacnaaraengtngtnathlaytnt 1202
Db	386 AGCCATACCTTCGCAAAGCCACAGGGGCAGAGTGGCTGTGAGGCATAGGGAAACCCACTCTT 445
OY	1203 ncaytygycaayaytngaygtctmgncaygngttnaarngngaycaytytgngcnytmg 1262
Db	446 GCATTCAGGTGATCCACCAAGCTGTGAGACATAGGTCTCAAGAGATVCAATTTTGGAGCTTTAAG 505

Oy	1263	ntcygagcygcnaengntlygmnaenctayatggnccnct-nccnytncygtcygnc	1321
Db	506	ATTGACACGCCCCGTCTGCATTTTCACAGACTTGTAAGGGCCAGTACGCCCTTTCTTTGGCC	565
Oy	1322	artcytcntlygnaengcngfntlyacnccartcytnttayttnccaytcygtat	1376
Db	566	AATTCTCCCATTTGGATGAGTGCTGTAATTTACCACCAATGCGCTCATTCATCTGTAT	620
RESULT	15		
LOCUS	AG097258	677 bp	DNA linear GSS 03-NOV-2001
DEFINITION	Pan troglodytes DNA, clone: PTB-098001.R, genomic survey sequence.		
ACCESSION	AG097258		
VERSION	AG097258.1	GI:16717775	
KEYWORDS	GSS; GSS (genome survey sequence).		
SOURCE	Pan troglodytes male lymphoblast DNA, clone_lib:PTB Chimpanzee Male BAC library clone:PTB-098001.R.		
ORGANISM	Pan troglodytes		
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Pan.		
AUTHORS	1 (sites)		
TITLE	Fujiyama, A., Hattori, M., Toyoda, A., Taylor, T. D., Yada, T., Tokoki, Y., Watanabe, H. and Sakaki, Y.		
REFERENCE	BAC end sequences of library PTB		
AUTHORS	Unpublished		
TITLE	2 (bases 1 to 677)		
JOURNAL	Fujiyama, A., Hattori, M., Toyoda, A., Taylor, T. D., Yada, T., Tokoki, Y., Watanabe, H. and Sakaki, Y.		
COMMENT	Submitted (02-AUG-2001) Asao Fujiyama, The Institute of Physical and Chemical Research (RIKEN), Genomic Sciences Center (GSC); 1-7-22 Shuhei-chou, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan (E-mail:shimbesegsc.riken.go.jp, URL:http://hgp.gsc.riken.go.jp/, Tel:81-45-503-9111, Fax:81-45-503-9170)		
FEATURES	Clones are derived from the chimpanzee BAC library PTB This BAC end was generated during the R&D process and may have higher chance of clone tracking errors.		
PRIMERS			
LIBRARY	Sequencing: M13Rev		
VECTOR	1 : pKS145		
R.Site 1	: SacI		
R.Site 2	: SacI		
Location/Qualifiers			
1..677			
/organism="Pan troglodytes"			
/db_xref="taxon:9598"			
/clone="PTB-098001.R"			
/sex="male"			
/cell_type="lymphoblast"			
/clone_lib="PTB Chimpanzee Male BAC library"			
BASE COUNT	178 a 175 c 176 g 147 t		1 others
ORIGIN			
Query Match	20.6%; Score 283.2; DB 12; Length 677;		
Best Local Similarity	48.8%; Pred. No. 1e-53;		
Matches 294; Conservative 113; Mismatches 187; Indels 9; Gaps 3;			
Oy	760	gongtngcngnagrtgngcngmncnnaarctnggcarytncnmsngngtngarcng	819
Db	68	GCTGGGCTTGAAGAAGGAGAAAGCCCAAGCCTTGCGACGCTTCATGTGGGTGAGGCT	127
Oy	820	gtungngcnaaraarwsmgnathargtntcyggarcncnccnathmgntlycaraarath	879
Db	128	ACAGGTGCAGAGAGATCAACAATGTAGAGTTGTGTGAACCTCACCTAGATT-AGACGATG	186
Oy	880	taygnaagccntcygatgctcmgmncaraarcttgcngtngngtngcngnmsntgtgmgn	939
Db	187	TATGAAACACACCTGGATGCCAGGATTAAGATTGCTGTCAGGGTGAAGGCCATCATGTGAGA	246

corresponding to the overlapping clone, as we submit sequences with only a small overlap as described above.

This sequence has been finished according to sequence map criteria as follows. An attempt is made to resolve all sequencing problems, such as compressions and repeats, but not necessarily within known annotated human repeat sequence elements (e.g. Alu) where the sequence is ambiguous, there is an annotation using the 'unsure' feature key.

The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases:

Em.: EMBL; SW: SWISSPROT; Tr.: TREMBL; Wp.: WormPEP; Information

on the WormPEP database can be found at

http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence was generated from part of bacterial clone contigs of human chromosome 22, constructed by the Sanger Centre Chromosome 22 Mapping Group. Further information can be found at

<http://www.sanger.ac.uk/HGP/Chr22>

RP1-302D9 is from the library RP1-1 constructed at the Roswell Park Cancer Institute by the group of Pieter de Jong. For further details see <http://bacpac.med.buffalo.edu/>

VECTOR: pCYPAC2

This sequence is the entire insert of clone RP1-302D9 the true left end of clone CTA-282F2 is at 69682 in this sequence. The true right end of clone CTA-415G2 is at 55167 in this sequence.

FEATURES

SOURCE

Location/Qualifiers

1..145880

/organism="Homo sapiens"

/db_xref="taxon:9606"

/chromosome="22"

/clone="RP1-302D9"

/clone_1fb="RP1-1"

188..245

/note="MER3 repeat: matches 144..209 of consensus"

246..571

/note="AluX repeat: matches 1..312 of consensus"

572..759

/note="MER3 repeat: matches 1..144 of consensus"

783..933

/note="MER3A repeat: matches 26..187 of consensus"

1033..1336

/note="AluSP repeat: matches 1..299 of consensus"

1450..1583

/note="MIR repeat: matches 24..160 of consensus"

1687..1752

/note="L2 repeat: matches 2593..2661 of consensus"

2350..2660

/note="AluSc repeat: matches 3..309 of consensus"

2684..2981

/note="AluSg repeat: matches 2..300 of consensus"

3323..3343

/note="MLTIE repeat: matches 116..136 of consensus"

3344..3652

/note="AluY repeat: matches 1..309 of consensus"

3653..3928

/note="MLTIE repeat: matches 136..359 of consensus"

3929..4278

/note="THERB repeat: matches 3..364 of consensus"

4279..4485

/note="MLTIE repeat: matches 359..568 of consensus"

5073..5176

/note="52 copies 2 mer ct 78 conserved"

5181..5491

/note="AluIb repeat: matches 1..311 of consensus"

6369..6485

/note="L2 repeat: matches 2579..2705 of consensus"

6647..6685

/note="MADL repeat: matches 1..23 of consensus"

6686..6987

/note="AluX repeat: matches 1..302 of consensus"

6988..7036

/note="MADL repeat: matches 23..77 of consensus"

7482..7754

/note="AluIb repeat: matches 9..290 of consensus"

repeat_region

7775..8060

/note="AluIb repeat: matches 1..295 of consensus"

repeat_region

8414..8551

/note="L2 repeat: matches 2553..2706 of consensus"

repeat_region

8914..9030

/note="MIR repeat: matches 147..262 of consensus"

repeat_region

9110..9280

/note="MIR repeat: matches 91..262 of consensus"

repeat_region

9283..9412

/note="MIR repeat: matches 15..144 of consensus"

repeat_region

9521..9679

/note="FAM repeat: matches 3..161 of consensus"

repeat_region

9820..10225

/note="MSTB repeat: matches 2..425 of consensus"

misc_feature

complement(10179..10678)

/note="match: GSS: Em:B56592"

misc_feature

complement(10204..10728)

/note="match: GSS: Em:A0701486"

misc_feature

complement(10249..10706)

/note="match: GSS: Em:A0225495"

repeat_region

10312..10383

/note="MIR repeat: matches 79..150 of consensus"

misc_feature

10718..11310

/note="match: GSS: Em:B14024"

misc_feature

10784..11201

/note="match: GSS: Em:B43656"

repeat_region

11838..11946

/note="MIR repeat: matches 20..137 of consensus"

repeat_region

12174..12445

/note="L2 repeat: matches 1988..2275 of consensus"

repeat_region

12444..12642

/note="MIR repeat: matches 63..241 of consensus"

misc_feature

13017..13369

/note="match: STS: Em:G49301"

repeat_region

13331..13397

/note="MIR repeat: matches 174..244 of consensus"

repeat_region

13398..13698

/note="AluSP repeat: matches 1..302 of consensus"

repeat_region

13699..13810

/note="MIR repeat: matches 76..174 of consensus"

repeat_region

13806..13919

/note="MIR repeat: matches 77..189 of consensus"

repeat_region

13945..14060

/note="MIR repeat: matches 24..142 of consensus"

repeat_region

14061..14367

/note="AluY repeat: matches 1..301 of consensus"

repeat_region

14368..14452

/note="MIR repeat: matches 141..225 of consensus"

repeat_region

14589..14679

/note="MIR repeat: matches 173..262 of consensus"

misc_feature

14597..15201

/note="match: GSS: Em:AQ553482"

misc_feature

14616..15060

/note="match: GSS: Em:AQ370601"

repeat_region

14868..15040

/note="MIR repeat: matches 49..233 of consensus"

repeat_region

15071..15188

/note="L2 repeat: matches 2112..2239 of consensus"

repeat_region

15304..15399

/note="MLTIB repeat: matches 1..99 of consensus"

repeat_region

15490..15662

/note="AluSg1 repeat: matches 2..114 of consensus"

repeat_region

15669..15727

/note="MLTIB repeat: matches 119..178 of consensus"

repeat_region

15728..16027

/note="AluSc repeat: matches 1..299 of consensus"

repeat_region

16028..16245

/note="MLTIB repeat: matches 178..390 of consensus"

repeat_region

16546..16854

/note="AluY repeat: matches 1..300 of consensus"

repeat_region

18296..18323

/note="MSTB repeat: matches 2..29 of consensus"

repeat_region

18324..18392

/note="MER66-internal repeat: matches 4919. .4993 of consensus"
repeat_region 18393. .18712
/note="AluJb repeat: matches 1. .311 of consensus"
repeat_region 18713. .19133
/note="MER66-internal repeat: matches 4548. .4919 of consensus"
complement(18872. .19230)
/note="match: GSS: Em:AQ005063"
19251. .19719
/note="match: GSS: Em:B14179"
19537. .20290
/note="HERVH21 repeat: matches 4657. .5784 of consensus"
repeat_region 20317. .20382
/note="33 copies 2 mer ta 68 conserved"
repeat_region 20513. .20666
/note="77 copies 2 mer tt 70 conserved"
repeat_region 20682. .21008
/note="AluSg1 repeat: matches 1. .306 of consensus"
repeat_region 21239. .21553
/note="HUR5-P3 repeat: matches 4410. .4713 of consensus"
repeat_region 21882. .22254
/note="THE1B repeat: matches 1. .364 of consensus"
repeat_region 22302. .22537
/note="MER66-internal repeat: matches 2186. .2417 of consensus"
repeat_region 22538. .22850
/note="AluSP repeat: matches 1. .313 of consensus"
repeat_region 22851. .23801
/note="MER66-internal repeat: matches 1210. .2186 of consensus"
repeat_region 23905. .23989
/note="MER66-internal repeat: matches 3017. .3102 of consensus"

Query Match 53.28; Score 921; DB 9; Length 145860;
Best Local Similarity 58.2%; Pred. No. 7.9e-215;
Matches 750; Conservative 300; Mismatches 239; Indels 0; Gaps 0;

Qy 443 tngnccnccgngngcncgngnynincncartaycngngarathsnaragacna 502
Db 37254 TGGAAATGCTCGTGCCGCCAGGTTTACACAAATATACAGGAATTAAGTAATGACAA 37313
Qy 503 artgycnltgycngayatharmngnswngntlyacngtlnaarytngnngnaarytnc 562
Db 37314 AATGCCCTGTCCTGATATAGAAAGTCAGCCTTACTGTGAAGTCAGTGAATAATTC 37373
Qy 563 cnytnccntlyaacrcnathlthtlyacngngntlyntayaygcncarmngaytlna 622
Db 37374 CTCTTCCTTCAAGCCCTCATCTTCAACAGGCGCTGTACAAATGCCAGAGGATTTAA 37433
Qy 623 argargcnatgngntlycngtngngnccngngnaaytlayaywsnswntlyagyg 682
Db 37434 AGGAGGCGCATGGAGTCTTCTTCAGGCGCTGGGAATTAATCAATTCAGCTTGATG 37493
Qy 683 tngartncaycaytlyaaargtlnaayathltyatnatmgnaarcarathyltngcnaaya 742
Db 37494 TTGAGCTCATCATTCGCAAGATGATATTTGGCTAATGACGAACAATTTTGGCTAATA 37553
Qy 743 argargarathwnaarcarcarwsnathlthtlyacngngntlyntayaygcncarmngaytlna 802
Db 37554 AGGAGGAATTTCTTAAGCAGCAAGCAATTCAGAGTGTGCTGTTAAAGGCAT 37613
Qy 803 tyantltatmgngargcngarcararwsnswngaraaytlncaaycngayaytlna 862
Db 37614 TCAAGTTATTAAGGAGGAGCAAGCATTAAGAGTCAAGAAATTTGACCCCTGCAATGTGA 37673
Qy 863 thaaaraaraaycncntlycngargnaartlyaaaytlnccngcngarathltyathl 922
Db 37674 TAAAAAAGAAAAACCCATTTTCTGAGGGGAATTCACACTGGCTGCCAACAATTTTCATAT 37733
Qy 923 gyaaygararalytlnaaytlnaaycncargayaygngargaraayathltsntgacntlyc 982
Db 37734 GTAATGAGAGCTGAATGTATCTCTCAAGACAAATGGGGAATAATATCTCTGAGCATGTG 37793

Qy 983 armgnwnswncarcarwsnathaarwnytnngcngtngmngcngmngnaartggtlyt 1042
Db 37794 AGAGTCTTCACAGCAGATTCATCAATCATCTGCTGGAGGCTGAGGAAAAATGTTT 37853
Qy 1043 yggngacngncngcngnswntlytlygtlytncarcngmngaytlytngtncntlytnc 1102
Db 37854 GTGGACAGGCCCAAGGCTCCCTGTCTGTGTGCACCTTACAGACTTGTGTGCTGTCC 37913
Qy 1103 cngtlnaaywsngcngtngcngsngargngcngswncncnaarctntgycarytncnswng 1162
Db 37914 CAGTTAATTCAGCTGTGCTTCAGAGGCTGCAAGGCCCAAGCCTTGGAGCTTCCAAAGTG 37973
Qy 1163 gngtngarcngtngngcnaaararwsnwnathltyatngtlyggarcncnctnmt 1222
Db 37974 GTGTTGACCTGTGGTGTCAAGAAAGTCAAGATTTGAGATTGGGAACTTCCATCAAT 38033
Qy 1223 tycaraarathlaygnaaycngtlygatycngmngcaratlytngcngtngngtngnw 1282
Db 38034 TTCAAGAGATATATGGAACCCCTGATGCGCAGCAGAGATTTGCTGAGGGGTGGGCT 38093
Qy 1283 snwsntlygmngacnswngcngmngtngtncaraargnaaytlytngtlyggarcncnc 1342
Db 38094 CCTCATGAGAAACCTCTGCAAGGCTACTACAAAGGAATGTTGGGTGGAGGCCCCAC 38153
Qy 1343 aymngtncnswngngcncnswngnswngngngtngmngnswncncnswngsm 1402
Db 38154 ACAGAGTCCCAAGTGCGCTCCATCTAGTAGAGCTGTAAGAAAGTCCACCATCTCCA 38213
Qy 1403 gnytcaraargngmngnswncnswngaytlncaaytngtncngaraararwsnagaya 1462
Db 38214 GACCTCAGAGGCTATATGGAACCCCTGATGCGCAGCAGATGTGCTGTAAGAAATTCACAGACA 38273
Qy 1463 cncartlycarcngtlnaaycngcngcngatvgararwsngtncntlayaarcngtngtng 1522
Db 38274 CTCACTGCGAGCTGTGAAAGCAGCAGGATGAGTCTGTACCTCAAAACCCGATGTGG 38333
Qy 1523 cngargtlnaarcngtngnathltyatngtlytncaytlycayaytlnaaygtngmng 1582
Db 38334 CAGAGCTGACCAAGACCGTGGGAATCTACCTGTGATTCATTCATGACCTGGAGCGGAAC 38393
Qy 1583 aygngtlnaarmngnayaaytlytngcngntlytngtlytngtlytngtlytngtlytng 1642
Db 38394 ATGAGCTCAAAAGAGATATTTTGGAGCTTAAAGATTTGACGTGCCCATCGATTTCCGA 38453
Qy 1643 cntlaytngcngtncnynlytngtlytngcncartlytlycncntlytngcngcngtnt 1702
Db 38454 CTATATGAGGCGCCCTACCCCTTGTGTTGGCCATTTTTCATTTGGAACTGCGCTAT 38513
Qy 1703 tyancartlytlytlnaaytlytngtlytngtlytngtlytngtlytngtlytng 1731
Db 38514 TTACCAATGCTGTACCTCCATTTGTATG 38542

RESULT 2
AC025577 154090 bp DNA linear PRI 25-AUG-2000
LOCUS Homo sapiens 12 BAC RP11-13C3 (Roswell Park Cancer Institute Human
DEFINITION BAC Library) complete sequence.
AC025577
VERSION AC025577.15 GI:9910028
KEYWORDS hng.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 154090)
Muzny D.M., Adams C., Adio-Oduola B., Ali-Osman F.R., Allen C.,
Aisbrooks S.L., Amaratunge H.C., Are J.R., Banks T., Barbataia J.,
Benton J., Bimaga K., Blankenburg K., Bonnin D., Bouck J.,
Bowle S., Brieval M., Brown E., Brown M., Bryant N.P., Buhaey C.,
Burch P., Burkett C., Burrell K.L., Byrd N.C., Caron T.F.,
Carter M., Cavazos S.R., Chacko J., Chavez D., Chen G., Chen R.,

Chen, Z., Chiu, D., Chowdhry, I., Christopoulos, C., Cleveland, C. D., Cox, C., Coyle, M. D., Dathorne, S. R., David, R., Davila, M. L., Davis, C., Davy, Carroll, L., Dederich, D. A., Delaney, K. R., Delgado, O., Denn, A. L., Ding, Y., Dinh, H. H., Douthwaite, K. J., Draper, H., Dugan-Rocha, S., Durbin, K. J., Earnhart, C., Edgar, D., Edwards, C. C., Elhaj, C., Emerling, S., Escotto, M., Falls, T., Ferraguto, D., Flagg, N., Ford, J., Foster, P., Frantz, P., Gabisi, A., Gao, J., Garcia, A., Garner, T., Garza, N., Gill, R., Gorelli, J. H., Guevara, W., Gunaratne, P., Hale, S., Hamilton, K., Han, J., Harris, C., Harris, K., Hart, M., Havlak, P., Hawes, A., Hernandez, J., Hernandez, O., Hodgson, A., Hoques, M., Hollway, C., Hollins, B., Homai, F., Howard, S., Huber, J., Huik, S., Hume, J., Ioshikhes, I., Jackson, L. E., Jacobson, B., Jia, Y., Johnson, R., Jolivet, S., Joudah, S., Karlsson, E., Kelly, S., Khan, U., King, L., Korvah, J., Kovar, C., Kratovic, J., Kureshi, A., Landry, N., Leal, B., Lee, E., Lewis, L. C., Lewis, L., Li, J., Li, Z., Licharge, O., Lieu, C., Liu, J., Liu, W., Lousegh, H., Lozano, R. J., Lu, X., Lucier, A., Lucier, R., Luna, R., Ma, J., Maheshwari, M., Mapa, P., Marondel, I., Martin, R., Martindale, A., Martinez, E., Massey, E., Mawhinney, E., McLeod, M. P., Meador, M., Mel, G., Merscher, S., Metzger, M., Miller, A., Miner, G., Miner, Z., Mitchell, T., Mohabbat, K., Montgomery, K. T., Morgan, M., Morris, S., Moser, M., Muzny, D., Neal, D., Nelson, D., Newton, J., Newton, N., Nguyen, A., Nguyen, N., Nguyen, N., Nickerson, E., Nwokoko, S., Oguh, M., Okwuonu, G., Oragunye, N., Oviedo, R., Pace, A., Payton, B., Peery, J., Perez, L., Peters, L., Pickens, R., Primus, E., Pu, L., Quiles, M., Ren, Y., Rivas, M., Rojas, A., Rojudo, K., Rolfe, M., Ruiz, S., Saverly, G., Scherer, S., Scott, G., Shen, H., Shum, C., Shooshari, N., Sisson, I., Sodergren, E., Sonalike, T., Sparks, A., Stanley, H., Stone, H., Sutton, A., Svatek, A., Taber, P., Tamerisa, K., Tang, H., Tansey, J., Taylor, C., Taylor, T., Tefford, B., Thomas, N., Thomas, S., Usmani, K., Vasquez, L., Vera, Y., Villalon, D., Vinson, R., Wall, R., Wang, S., Ward-Moore, S., Warren, R., Washington, C., Wallington, S., Williams, G., Williamson, A., Wleczek, R., Woodem, S., Worley, K., Wu, C., Wu, Y., Wu, Y. F., Zhou, J., Zorrilla, S., Kucherlapati, R., Nelson, D. and Gibbs, R.

Direct Submission
Unpublished
2 (bases 1 to 154090)
Worley, K. C.
Direct Submission
Submitted (11-MAR-2000) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 154090)
Worley, K. C.
Direct Submission
Submitted (25-AUG-2000) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
On Aug 25, 2000 this sequence version replaced gi:9664948.
INFORMATION: <http://www.hgsc.bcm.tmc.edu/> or email gc-help@bcm.tmc.edu

CLONE LENGTH: This sequence does not necessarily represent the entire insert of this clone. Overlapping regions of clones are only sequenced and submitted once, so the sequence for the remainder of the insert may be found in the record for the adjacent clones. Overlapping clones are noted at the beginning and end of the Features listing.

ANNOTATION OF FEATURES:
STSs are identified using EPCR (Genome Res. 7:541-550) searches of a local database that includes entries from dbSTS, GDB, and local mapping efforts.

Repeats are identified using RepeatMasker (A. Smit and P. Green, unpublished.) for Human and Mouse sequences.

Genes and Region of sequence similarity are identified by BLAST (Nuc. Acids Res. 25:3383-3402) similarity (expect < 1e-34) to the EST and cDNA sequences. Genes demonstrate at least two exons flanked by consensus splice sites that maintained sequence continuity across the splice junctions. Sequences that are not identical matches are annotated as similar.

SEQUENCING READ COVERAGE: Sequencing is completed to a minimum standard of double strand coverage with a minimum of 2 clones and 2 reads with no ambiguities or 2 chemistries with a minimum of 2 clones and 3 reads with no ambiguities. If the sequence quality for a region does not meet this standard, it will be indicated in the annotation as Low Coverage.

QUALITY OF INDIVIDUAL BASES: This sequence meets stringent quality standards - estimated error rate less than 1 per 10,000 bases. Reports of lowest quality individual bases and measures of base quality are listed below. Description of the metrics can be found at URL: <http://gc.bcm.tmc.edu:8088/quality.info/genbank.annotation.html>.

QUALSTAT-REPORT

----- Summary Statistics -----
Contig length: 154090
Phrap values in estimate: 15345
Average error rate (BCM-Phrap estimate): 0.000261755
Fraction of Phrap values less than 40 : 0.0284086
Number of consensus changing edits: 22
Number of N's in consensus : 0

Position	Original/Context	Consensus changing edits	Edited/Context
9347	aacacagacc(n)tttttttt	aacacagacc(t)tttttttt	aacacagacc(t)tttttttt
10135	attggccct(n)taagaaga	attggccct(t)taagaaga	attggccct(t)taagaaga
10353	ctgtccacc(n)aaagatcca	ctgtccacc(t)aaagatcca	ctgtccacc(t)aaagatcca
10417	agcaatgcag(n)gctacagaa	agcaatgcag(t)gctacagaa	agcaatgcag(t)gctacagaa
61959	cttactact(n)tggttatct	cttactact(t)tggttatct	cttactact(t)tggttatct
61960	tttaactatn(t)gttatctca	tttaactatn(t)gttatctca	tttaactatn(t)gttatctca
61961	ttaactatn(t)gttatctca	ttaactatn(t)gttatctca	ttaactatn(t)gttatctca
61982	acaacacag(n)taggttttg	acaacacag(t)taggttttg	acaacacag(t)taggttttg
61983	caacacag(n)taggttttg	caacacag(t)taggttttg	caacacag(t)taggttttg
61993	ttaggttttg(n)ttactcttc	ttaggttttg(t)ttactcttc	ttaggttttg(t)ttactcttc
62305	tcattgacc(n)ctgtacatc	tcattgacc(t)ctgtacatc	tcattgacc(t)ctgtacatc
62356	actgcaacc(n)tgctccacg	actgcaacc(t)tgctccacg	actgcaacc(t)tgctccacg
63722	ttacacata(n)cgatcacga	ttacacata(t)cgatcacga	ttacacata(t)cgatcacga
90212	cctagaanaa(n)gacttttct	cctagaanaa(t)gacttttct	cctagaanaa(t)gacttttct
90216	gaaanaagac(n)ttttcttct	gaaanaagac(t)ttttcttct	gaaanaagac(t)ttttcttct
90238	ttcttttta(n)ttttcttat	ttcttttta(t)ttttcttat	ttcttttta(t)ttttcttat
90236	ttttttttc(n)atcttaggg	ttttttttc(t)atcttaggg	ttttttttc(t)atcttaggg
90443	accttagag(n)ctctccacg	accttagag(t)ctctccacg	accttagag(t)ctctccacg
91405	caacagcttaa(n)caaggtaga	caacagcttaa(t)caaggtaga	caacagcttaa(t)caaggtaga
122290	aaacaaagaa(n)gtaagagaaa	aaacaaagaa(t)gtaagagaaa	aaacaaagaa(t)gtaagagaaa
137090	acagagaag(n)caaacacac	acagagaag(t)caaacacac	acagagaag(t)caaacacac
141632	tgaagccact(n)gaaagtaat	tgaagccact(t)gaaagtaat	tgaagccact(t)gaaagtaat

----- Distribution of Quality < 40 Bases -----

#	1000	900	800	700	600	500	400	300	200	100	0
bases	1000	900	800	700	600	500	400	300	200	100	0
Phrap Value Range	5	10	15	20	25	30	35	40			

FEATURES

----- Version: 1.01 gxf. -----
Location/Qualifiers

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source
1. .154090
/organism="Homo sapiens"
/db.xref="taxon:9606"
/chromosome="12"
/clone="RP11-13C3"
54. .288
/rpl_family="MIR"
repeat_region
1020. .1146
/rpl_family="MER3"
complement(1248. .1584)
repeat_region
/rpl_family="MER4B"
complement(1624. .1682)
repeat_region
/rpl_family="MIR"
2630. .2761
/rpl_family="GA-rich"
repeat_region
4088. .4216
/rpl_family="L2"
repeat_region
5232. .5301
/rpl_family="MER102"
complement(5450. .5749)
repeat_region
/rpl_family="AluY"
6571. .6616
/rpl_family="TGn"
repeat_region
7419. .7454
/rpl_family="L1ME3"
complement(7741. .7807)
repeat_region
8113. .8267
/rpl_family="MER5A"
repeat_region
8385. .8587
/rpl_family="MER5A"
repeat_region
8589. .8630
/rpl_family="MIR"
repeat_region
8666. .8695
/rpl_family="L2"
repeat_region
/rpl_family="(TFA)n"

Query Match 28.88; Score 497.8; DB 9; Length 154090;
Best Local Similarity 47.78; Pred. No. 1.6e-110;
Matches 488; Conservative 216; Mismatches 282; Indels 36; Gaps 2;
```

```
Qy 1189 wsmnqatthgargtntgggarcnccnathmgntlycaaraatltaggnaaycncntg 1248
:::|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 102113 TCAGAAATTTGAGGCTTGGGAGGCTTCACCTAGATTTCAGAGAGATTATAGGAATCCCTGG 102172
Qy 1249 atgcnmgnacaaatlygcnngtngngtngngnswnsntgmgnaacnswngcnmgntn 1308
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 102173 ATGTCCAGGAGAAAGTTTGTTCAGGGGTGGGCCCTCATGAGAAACCTCTGTGGGCA 102232
Qy 1309 gtncaraargnaaygtngntggtgarcncncncaymgntncnswngcnmgcnswn 1368
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 102233 GTGCGAAGGGAATGTGGGTGGGCTTGGAGCCATACAGAGATCCCTACCTGCGGGCACCACT 102292
Qy 1369 wsmnngcnngtngmgmgswncnccnswnsnsmngntncncaargmgmgswnaagay 1428
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 102293 AGGGAGCTGTGAGAAAGGGCCACCATCTCTCCACAGCCCGAGAAATGTAGTACCAAC 102352
Qy 1429 wsnlyncarcaygtncncgnaaraarwnacnagayacnartlycarrcngtlnaargcncn 1488
:::|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 102353 AGCTTGACACCATGTACTGGAAGAACACACAGAGACTCAACACAGCCCATGAAACAGACC 102412
Qy 1489 ggnatggarwngtncntctayaaaracngtngtngcngarytnacnaaracngtngnaeth 1548
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 102413 AAGAGGGAGCCTGTACCTCGAAGAACACAGAGAGAGCTTCCAGACCATGGGAAC 102472
Qy 1549 laytlnyrcaytlycaygaytngaygtmgncaygngtlnaarmgngaycaytlyggn 1608
:::|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 102473 CACCTCTTGCACTAGCTGAGCTAGATGTGAACATGCAAGTCAAGAGATCATATTGA 102532
Qy 1609 gcnylmngntlygaytlygcnaacngntlymgnaentlayatgngncngnt-nccnlyntg 1667
:::|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 102533 ACTTGGAGATTGTGACTACCTCTCTGCGTTTGCAGACTGCAGTGGGCGCATATAGCCCTTTG 102592
Qy 1668 yltvgncartlytlycncntlygngnagcngntlytncncartlytntaytlncaytg 1727
:::|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 102593 TTTTGCCCAATTTCCCATTTAGAACAGCTGTATTACTCAATGCCGTATGCCCATTTG 102652
Qy 1728 ya 1729
:::|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 102653 CA 102654

RESULT 3
AC026107 186660 bp DNA linear HTG 06-FEB-2002
LOCUS Homo sapiens chromosome 12 clone RP11-307L1, WORKING DRAFT
DEFINITION
SEQUENCE, 1 unordered pieces.
ACCESSION AC026107
VERSION AC026107.22 GI:13899178
KEYWORDS HTG; HTGS_PHASE1; HTGS_DRAFT; HTGS_FULLTOP.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 186660)
Munzy D.M., Adams C., Adio-Oduola B., Ali-oman F.R., Allen C.,
Aldbrooks S.L., Amaralunge H.C., Are J.R., Ayale M., Banks T.,
Bardella J., Benton J., Bimaye K., Blamendour K., Bonnin D.,
Bouck J., Bowie S., Brieva M., Brown E., Brown M., Bryant N.P.,
Buhay C., Burck P., Burkett C., Burrell K.L., Byrd N.C.,
Carroll T.F., Carter M., Cavazos S.R., Chacho J., Chavez D.,
Chen G., Chen R., Chen Z., Chowdhury I., Christopoulos C.,
Cleveland C.D., Cox C., Coyle M.D., Dethorne S.R., David R.,
Davila M.L., Davis C., Davy-Carroll L., Dederich D.A.,
Delaney K.R., Delgado O., Denn A.L., Ding Y., Dinh H.H.,
Douthwaite K.J., Draper H., Dugan-Rocha S., Durbin K.H.,
Earnhart C., Edgar D., Edwards C.C., Elhaj C., Escotto M.,
Falls T., Ferraguto D., Flagg N., Ford J., Foster P., Frantz P.,
Gabisi A., Geo J., Garcia A., Garner T., Garza N., Gill R.,
Gorrell J., Guera W., Gunaratne P., Hale S., Hamilton K.,
Harris C., Harris R., Hart M., Havlik P., Hawes A., Hernandez J.,
Hernandez O., Hodgson A., Hogues M., Holloway C., Hollins B.,
Honsi F., Howard S., Huber J., Huliy S., Hume J., Jackson L.E.,
```

Jacobson, B., Jia, Y., Johnson, R., Jolivet, S., Joudah, S.,
 Karlsson, E., Kelly, S., Khan, U., King, L., Korrah, J., Kovar, C.,
 Kratovic, J., Kuresh, A., Landry, N., Leal, B., Lewis, L.C., Lewis, L.,
 Li, J., Li, Z., Lichtarge, O., Lien, C., Liu, J., Liu, W., Louisgeed, H.,
 Lozadó, R., Lu, X., Lucier, A., Luckner, R., Luna, R., Ma, J.,
 Mesheshwari, M., Mapua, P., Martin, R., Martindale, A., Martinez, E.,
 Messer, E., Mawhinney, E., McLeod, M.P., Meador, M., Mel, G., Metzger, M.,
 Miner, G., Miner, Z., Mitchell, T., Mohabbat, K., Morgan, M., Morris, S.,
 Moser, M., Neal, D., Newton, J., Newton, N., Nguyen, A., Nguyen, N.,
 Nguyen, N., Nickerson, E., Nwokwenkwo, S., Ogutu, M., Okunolu, G.,
 Otágujare, N., Oviedo, R., Pace, A., Payton, B., Peery, J., Perez, L.,
 Peters, L., Pickens, R., Primus, E., Pu, L., Qulles, M., Ren, Y.,
 Rives, M., Rojas, A., Rojibokan, I., Rolfe, M., Ruiz, S., Savery, G.,
 Scherer, S., Scott, G., Shen, H., Shooshitari, N., Sisson, I.,
 Sodergren, E., Sonalke, T., Sparks, A., Stanley, H., Stone, H.,
 Sutton, A., Svatek, A., Tabor, P., Tameris, A., Tameris, K., Tang, H.,
 Tanton, J., Taylor, C., Taylor, T., Telford, B., Thomas, N., Thomas, S.,
 Usmani, K., Vasquez, L., Vera, V., Villalón, D., Vinson, R., Wall, R.,
 Wang, S., Ward-Moore, S., Warren, R., Washington, C., Wallington, S.,
 Williams, G., Williamson, A., Wleceyk, R., Woodson, S., Worley, K.,
 Wu, C., Wu, Y., Wu, Y. F., Zhou, J., Zorrilla, S., Nelson, D.,
 Weinstock, G. and Gibbs, R.
 Direct Submission
 Unpublished
 2 (bases 1 to 186660)
 Worley, K.C.
 Direct Submission
 Submitted (19-MAR-2000) Human Genome Sequencing Center, Department
 of Molecular and Human Genetics, Baylor College of Medicine, One
 Baylor Plaza, Houston, TX 77030, USA
 On May 1, 2001 this sequence version replaced gi:13871715.

```

* NOTE: Estimated insert size may differ from sequence length
* (see http://www.hgsc.bcm.tmc.edu/docs/genbank\_draft\_data.html)
*
* NOTE: This is a 'working draft' sequence. It currently
* consists of 1 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
*
*       1 186660: contig of 186660 bp in length.

```

FEATURES

BASE COUNT	57089 a	37346 c	35962 g	56263 t
ORIGIN				

[illegible]

LOCUS	AC008799	123070 bp	DNA	linear	HTG-18-JUL-2000
DEFINITION	Homo sapiens chromosome 5 clone CTD-2061E19, WORKING DRAFT SEQUENCE, 8 ordered pieces.				
ACCESSION	AC008799				
VERSION	AC008799.4	GI:9256046			
KEYWORDS	HTG; HTGS_PHASE2; HTGS_DRAFT.				
SOURCE	human.				
ORGANISM	Homo sapiens				
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.				
AUTHORS	1 (bases 1 to 123070)				
TITLE	Sequencing of Human Chromosome 5				
JOURNAL	Unpublished				
REFERENCE	2 (bases 1 to 123070)				
AUTHORS	DOE Joint Genome Institute.				
TITLE	Direct Submission				
JOURNAL	Submitted (03-ANG-1999) Production Sequencing Facility, DOE Joint Genome Institute, 2800 Mitchell Drive, Walnut Creek, CA 94598, USA				
COMMENT	On Jul 18, 2000 this sequence version replaced gi:7709316.				

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Web site: http://www.jgi.doe.gov
-----
Project Information
Center Project Name: 651088
Center clone name: CITB-H1_2061E19

```

Summary Statistics	
Consensus quality:	116295 bases at least Q40
consensus quality:	121288 bases at least Q30
Consensus quality:	122086 bases at least Q20
Estimated insert size:	123000, pulse field gel estimation
Estimated insert size:	122770, sum-of-contigs estimation
Quality coverage:	6.04 in Q20 bases; pulse field gel estimation
Quality coverage:	6.05 in Q20 bases; sum-of-contigs estimation

* NOTE: This is a 'working draft' sequence. It currently

* consists of n collages. Gaps between the collages
* are represented as runs of N . The order of the pieces
* is believed to be correct as given, however the sizes
* of the gaps between them are based on estimates that have
* provided by the submittor.

- * this sequence will be replaced
- * by the finished sequence as soon as it is available and
- * the accession number will be preserved.

*	9436:	contig of 9436 bp in length
*	9437	9536: gap of unknown length
*	9537	26537: contig of 17001 bp in length
*	26538	26637: gap of unknown length
*	26638	45958: contig of 19321 bp in length
*	45959	46058: gap of unknown length
*	46059	68657: contig of 22599 bp in length
*	68658	68757: gap of unknown length
*	68758	72152: contig of 3385 bp in length
*	72153	72252: gap of unknown length
*	72253	81069: contig of 8817 bp in length
*	81070	81169: gap of unknown length
*	81170	121547: contig of 40378 bp in length
*	121548	121644: gap of unknown length
*	121648	122070: contig of 1433 bp in length

FEATURES

Location/Qualifiers

```

SOURCE
1. :12307 "Homo sapiens"
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="5"
/clone="C7D-206IE19"
/clone_11p="caltech human BAC library D"
37763 a 24223 c 23855 g 36528 t 701 others
BASE COUNT
ORIGIN

```

Query Match 28.58; Score 493; DB 2; Length 123070;

Best Local Similarity 46.7%; Pred. No. 2.3e-109;
Matches 495; Conservative 220; Mismatches 309; Indels 36; Gaps 2;

Qy	706	aaytlhgytlnatgmnaarcaralhtylngcnaaayaatgaargaralhtsnarcarar	765
Db	107907	AACTTTAACTTGAAGAAATATGATTTAGGTAATCTGGCAAGAAATTTCTTAAGCCAA	107966
Qy	766	wsnathargarbnacnltgglnhylnythaaragcnltwstlyatlhmngargcnar	825
Db	107967	AGCATTTCCAGAGGCTTGGGCTGCTTTAAAGTATTCAGTTTAAAGGACAGAG	108026
Qy	826	cayaarwnsnrgnaaaylncayccngayaaygltnalhaararaaaycntllysn	885
Db	108027	CATPAAAGTCGGAATAATTTGCGAGTCGCAGACATGTGATAGAAAGAAATTCATTTCT	108086
Qy	886	garjgnaarltlyaarlyncngcnngarathtylathtyaaayaraarylnaaylnaay	945
Db	108087	GAGGAGAAATTTCAAGCTGGCTGCAGAAATATGATATGATACAGAGAGCTGAATATTAAT	108146
Qy	946	ccnargayaaygngnaraayalhtwsntlgactblycarmngsnwnsncarawnath	1005
Db	108147	CCCCAAGACAAATGGGGAAAAATCTCCAAAGCATGTCAAGAGCTTCACAGGACGCCCTC	108206
Qy	1006	aarvsnlyngcnltgmgncnmngmng--naarltgltlytyvgnaacngnccngnwns	1062
Db	108207	CCATCATTAACCCCAAAGGCTAGAGAGAAATAAGTTGTGTGGCCAGGCCAGGCTCC	108266
Qy	1063	ytlngytygynarccnmngnaryltngtncnltgytlnccnglnaa-----	1109
Db	108267	TTTGCTGTGTGACAGCTTAGGAGATTTGTGCCCTGCATCCACCTGCTCCAGCATAGCT	108326
Qy	1110	-----ywsnngcnltgcnwsnrgnargngcnwnsncnnaarcnltg	1149
Db	108327	GAAGGGGCCAAACTAGAGCTGGGGCTGTGGCTTCAAGAGTGCAGGCCCAAGCCTTG	108386
Qy	1150	carlytncnwnsnngnltngarcngltngnngcnaaraarwsnmgnalhtarylntvgar	1209
Db	108387	CAGCTTCATGTGCTGTGAGCCCTGCAGAGTGCACAGAAAGTCAACAAATTTGGGGTTTGGAA	108446
Qy	1210	cnccnnaalmngnttlycraaralhtlayjgnaaycmtlgatlgcmngnaraarltyn	1269
Db	108447	CCTCCACCTAATTTGCAGATGATATGAGAAATGCCCGGATGACCAAGCAAGATTGCT	108506
Qy	1270	gtngnngltngnwnsnwtgmgngacnwnsgcmngnltngtncaraargnaayltngn	1329
Db	108507	GCAGGGGACGGCCTTCAATGAGAACTCTGTAGGSCATGTGAGAAAGAAATGTAGGG	108566
Qy	1330	ltggarccnccncaymngnltcnwnsnngnccnwnsnwnsmngnngcnltmgmngwns	1389
Db	108567	TCAGAGCCCAACACAGAGTACCACATAGAGCACTGCTGTGGAGCCTGTGAGAAAGAG	108626
Qy	1390	cnccnwnsnwmngnlyncaraargmngmwnsnacngaysnlytnarcarylncngar	1449
Db	108627	CCACCATCTTCCAACTCCAAATGATAGGTCCCAACAGCTTGCACTGTACTGGA	108686
Qy	1450	aarwnacnayaacnartlycarcnngthaaragcnngnaltgawrsgnltncntay	1509
Db	108687	AAAGCCACAGGCACTCATTTGCCACCTGTGAACACACAGAGGAGAGCTATCCCTGC	108746
Qy	1510	aaaracnltngltngcnaryltnacnaaraacnltngnaltlayltnylncaytygcayay	1569
Db	108747	AAAGTCACAGGGGCGAAGCTGTCCAAAGACCTTGGGAACCTACCTTGCTCCTCAGCGTGAC	108806
Qy	1570	yltngaygtmngncaygngnltaaarmngayayltlytngnrcnyltmgntlygatyccn	1629
Db	108807	CCGGATGTGAGAACATGTGAGTAAAGGAGATCATATTTTGGACTTTAAGATTGTACTGCCCT	108866
Qy	1630	acnngnltymnacntayaltgngcnngtncnrcnyltngtytggncarltlyccntly	1689
Db	108867	GCTGATTTTCAGAGCTGCATGGGGCCTGTAGCCCTTTGCTTTGGCCCAATTTTCGCCATT	108926
Qy	1690	ggnacnngnltlyacnrcarlytlnylnaytncaytga 1729	

Db 108927 GGACAGCTGATTATACCCATGCTGACCCGATGTGA 108966

RESULT 6
AC044889/c 189768 bp DNA linear HTG 22-MAY-2000
LOCUS Homo sapiens chromosome 2 clone RP11-792C1 map 2, WORKING DRAFT
DEFINITION
SEQUENCE, 36 unordered pieces.
AC044889
AC044889.2 GI:8016676
KEYWORDS HTG; HTGS_PHASE1; HTGS_DRAFT.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE
1 (bases 1 to 189768)
Birten,B., Linton,L., Nusbaum,C. and Lander,E.
Homo sapiens chromosome 2, clone RP11-792C1
Unpublished
2 (bases 1 to 189768)
Birten,B., Linton,L., Nusbaum,C., Lander,E., Abraham,H., Allen,N.,
Anderson,S., Baldwin,J., Barna,N., Bastien,V., Beda,F.,
Boguslavsky,L., Bouhgalter,B., Brown,A., Burkett,G.,
Campoliano,A., Castle,A., Choepel,Y., Colangelo,M., Collins,S.,
Collamore,A., Cooke,P., Dearlano,K., Dewar,K., Diaz,J.S.,
Dodghe,S., Domino,M., Doyle,M., Ferreira,P., Fitzhugh,W., Gage,D.,
Galagan,J., Gardyna,S., Ginde,S., Goyette,M., Graham,L.,
Grand-Pierre,N., Grant,G., Hagos,B., Heatford,A., Horton,L.,
Klein,D., Laroque,K., Lamazares,R., Landers,T., Lehoczy,J.,
Levine,R., Liu,C., Liu,G., Locke,K., Macdonald,P., Marcus,N.,
McCarthy,M., McEwan,P., McGurk,A., McKernan,K., McPheters,R.,
Meldrum,J., Menkus,L., Mihova,T., Miranda,C., Mlenge,V., Morrow,J.,
Murphy,T., Naylor,J., Norman,C.H., O'Connor,T., O'Donnell,P.,
O'Neill,D., Oliver,T.M., Oliver,J., Peterson,K., Pierre,N.,
Pisan,C., Pollara,V., Raymond,C., Riley,R., Rogov,P., Rothman,D.,
Roy,A., Santos,R., Schauer,S., Severy,P., Spencer,B.,
Stange-Thomann,N., Stojanovic,N., Subramanian,A., Talamas,J.,
Teifage,S., Theodore,J., Tirrell,A., Travers,M., Triggillo,J.,
Vassiliev,H., Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W.J.,
Young,G., Zainoun,D., Zimmer,A. and Zody,M.

TITLE
JOURNAL
COMMENT
Direct Submission
Submitted (12-APR-2000) Whitehead Institute/MIT Center for Genome
Research, 320 Charles street, Cambridge, MA 02141, USA
On May 22, 2000 this sequence version replaced gi:7543856.
All repeats were identified using RepeatMasker:
Smt, A.F.A. & Green, P. (1996-1997)
http://ftp.genome.washington.edu/RM/RepeatMasker.html

Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WtBR

Web site: http://www-seq.wi.mit.edu

Contact: sequence_submissions@genome.wi.mit.edu

Project Information

Center project name: L9594

Center clone name: 792_C_1

Summary Statistics

Sequencing vector: M13, M7815, 100% of reads

Chemistry: Dye-terminator Big Dye, 100% of reads

Assembly program: Phrap; version 0.960731

Consensus quality: 168749 bases at least Q40

Consensus quality: 180047 bases at least Q30

Consensus quality: 184014 bases at least Q20

Insert size: 190000; agarose-fp

Insert size: 186266; sum-of-contigs

Quality coverage: 3.8 in Q20 bases; agarose-fp

Quality coverage: 3.9 in Q20 bases; sum-of-contigs

NOTE: This is a 'working draft' sequence. It currently
* consists of 36 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.

* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
1 1015: contig of 1015 bp in length
1016 1115: gap of 100 bp
1116 2284: contig of 1169 bp in length
2285 2384: gap of 100 bp
2385 3898: contig of 1514 bp in length
3899 5253: gap of 100 bp
5254 5359: contig of 1261 bp in length
5360 6912: gap of 1553 bp in length
6913 7012: gap of 100 bp
7013 8145: contig of 1133 bp in length
8146 8245: gap of 100 bp
8246 9694: contig of 1449 bp in length
9695 9794: gap of 100 bp
9795 11004: contig of 1210 bp in length
11005 11104: gap of 100 bp
11105 12907: contig of 1803 bp in length
12908 13007: gap of 100 bp
13008 14916: contig of 1909 bp in length
14917 15016: gap of 100 bp
15017 17114: contig of 2098 bp in length
17115 17214: gap of 100 bp
17215 18527: contig of 1313 bp in length
18528 18627: gap of 100 bp
18628 20561: contig of 1934 bp in length
20562 20661: gap of 100 bp
20662 22484: contig of 1823 bp in length
22485 22584: gap of 100 bp
22585 25287: contig of 2703 bp in length
25288 25387: gap of 100 bp
25388 27603: contig of 2216 bp in length
27604 27703: gap of 100 bp
27704 30020: contig of 2217 bp in length
30021 30120: gap of 100 bp
30121 33472: contig of 3352 bp in length
33473 33572: gap of 100 bp
33573 36084: contig of 4512 bp in length
36085 38184: gap of 100 bp
38185 41069: contig of 2885 bp in length
41070 41169: gap of 100 bp
41170 45266: contig of 4097 bp in length
45267 45366: gap of 100 bp
45367 51092: contig of 5726 bp in length
51093 51192: gap of 100 bp
51193 55262: contig of 4070 bp in length
55263 55362: gap of 100 bp
55363 60982: contig of 5620 bp in length
60983 61082: gap of 100 bp
61083 67628: contig of 6546 bp in length
67629 67728: gap of 100 bp
67729 72690: contig of 4962 bp in length
72691 72790: gap of 100 bp
72791 79286: contig of 6496 bp in length
79287 79386: gap of 100 bp
79387 83451: contig of 4065 bp in length
83452 83551: gap of 100 bp
83552 92170: contig of 8619 bp in length
92171 92270: gap of 100 bp
92271 96333: contig of 4063 bp in length
96334 96433: gap of 100 bp
96434 103218: contig of 6785 bp in length
103219 103318: gap of 100 bp
103319 112553: contig of 9235 bp in length
112554 112653: gap of 100 bp
112654 123239: contig of 10586 bp in length
123240 123339: gap of 100 bp
123340 137921: contig of 14582 bp in length
137922 138021: gap of 100 bp
138022 157913: contig of 19892 bp in length
157914 158013: gap of 100 bp

Db 84123 ATTTGCTGTATTATCCATGCCTGTACTGCCAATGTTT 84161

RESULT 8
AC025233/c
LOCUS
DEFINITION
AC025233 161575 bp DNA linear HTG 07-JUL-2000
Homo sapiens chromosome 17 clone RP11-333E1, WORKING DRAFT
SEQUENCE, 27 unordered pieces.
AC025233
AC025233.4 GI:8954317
KEYWORDS
HTG: HTGS_PHASE1; HTGS_DRAFT.
SOURCE
ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE
AUTHORS
TITLE
JOURNAL
REFERENCE
AUTHORS
TITLE
JOURNAL
Unpublished
2 (bases 1 to 161575)
Waterston, R.H.
Direct Submission
Submitted (07-MAR-2000) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
On Jul 7, 2000 this sequence version replaced gi:7801489.

COMMENT

----- Genome Center -----
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu/gsc/index.shtml
----- Project Information -----
Center project name: H_NH0335E01
----- Summary Statistics -----
Sequencing vector: M13; 90%
Sequencing vector: plasmid; 10%
Chemistry: Dye-primer ET; 81% of reads
Chemistry: Dye-terminator Big Dye; 19% of reads
Assembly program: Phrap; version 0.990319
Consensus quality: 144558 bases at least Q40
Consensus quality: 149743 bases at least Q30
Consensus quality: 152625 bases at least Q20
Insert size: 176000; agarose-fp
Insert size: 158975; sum-of-ctrls
Quality coverage: 3.48 in Q20 bases; agarose-fp
Quality coverage: 3.75 in Q20 bases; sum-of-ctrls

* NOTE: This is a 'working draft' sequence. It currently
* consists of 27 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
*
* 1 1141: contig of 1141 bp in length
* 1142 1241: gap of unknown length
* 1242 2424: contig of 1183 bp in length
* 2425 2524: gap of unknown length
* 2525 3961: contig of 1437 bp in length
* 3962 4061: gap of unknown length
* 4062 5712: contig of 1651 bp in length
* 5713 5812: gap of unknown length
* 5813 8019: contig of 2207 bp in length
* 8020 8120: gap of unknown length
* 8120 10208: contig of 2089 bp in length
* 10209 10309: gap of unknown length
* 10309 11734: contig of 1426 bp in length
* 11735 11834: gap of unknown length
* 11835 13265: contig of 1431 bp in length
* 13266 13366: gap of unknown length
* 13366 14425: contig of 1060 bp in length
* 14426 14525: gap of unknown length

* 14526 15609: contig of 1084 bp in length
* 15610 15709: gap of unknown length
* 15710 17529: contig of 1820 bp in length
* 17530 17629: gap of unknown length
* 17630 20351: contig of 2722 bp in length
* 20352 20451: gap of unknown length
* 20452 23753: contig of 3302 bp in length
* 23754 23853: gap of unknown length
* 23854 26537: contig of 2684 bp in length
* 26538 26637: gap of unknown length
* 26638 31339: contig of 4702 bp in length
* 31340 31439: gap of unknown length
* 31440 37041: contig of 5602 bp in length
* 37042 37141: gap of unknown length
* 37142 45268: contig of 8127 bp in length
* 45269 45368: gap of unknown length
* 45369 52323: contig of 6955 bp in length
* 52324 52423: gap of unknown length
* 52424 60239: contig of 7816 bp in length
* 60240 60339: gap of unknown length
* 60340 68385: contig of 8046 bp in length
* 68386 68485: gap of unknown length
* 68486 78834: contig of 10349 bp in length
* 78835 78934: gap of unknown length
* 78935 90422: contig of 11488 bp in length
* 90423 90522: gap of unknown length
* 90523 100763: contig of 10241 bp in length
* 100764 100863: gap of unknown length
* 100864 111672: contig of 10809 bp in length
* 111673 111772: gap of unknown length
* 111773 126855: contig of 15083 bp in length
* 126856 126955: gap of unknown length
* 126956 142557: contig of 15602 bp in length
* 142558 142657: gap of unknown length
* 142658 161575: contig of 18918 bp in length.
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Location/Qualifiers
1. 161575
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="RP11-333E1"

BASE COUNT 44288 a 35882 c 35114 g 43686 t 2605 others
ORIGIN

Query Match 28.2% Score 488.8; DB 2; Length 161575;
Best Local Similarity 48.5%; Pred. No. 2.8e-108;
Matches 485; Conservative 202; Mismatches 280; Indels 34; Gaps 2;

Qy 744 rgargarthhsnaarcarcarwanatbpcargatgtnacntgggtbnylnyaaagcntt 803
Db 121612 GGAAAGAAATTTCTAAGCGCAAAACATTCAAGGTGTGACTTGCGTAAATGGATT 121553

Qy 804 ywsntlyatlmngargngargncayaaarwsnsgaraaytlnaaycngayaaygtnat 863
Db 121552 CAGTTTATGAGGGAGAGAGAGCATTAAGAAATTTGCAGCCTGAGAAATGGAT 121493

Qy 864 haaraaraaraaycnclywsngargnaartlyaaaytngcngcngarathychtg 923
Db 121492 AGAAAGAAAGAAACCATTTCTGAGGAGAAATTCATCAGCTCAGAAATTTGCATPAG 121433

Qy 924 yaaygarayrtnaaygnayccncargayaaygngarayaalhsntgacntgyca 983
Db 121432 TAACAAGAGCGTAATGTTAATCCGCAAGACAAATGGGAAATGTCTCCAGGCAATGCA 121373

Qy 984 tmgwsnwsnrcarcarwsnatlaarwsnyltngcyltngcngcngmgngnaartgltctg 1043
Db 121372 GAAGTCTTCACAGAGCCGCCCTCCATCAGAGCCTGGAGGCGCTGAGAAATGTTTTC 121313

Qy 1044 ygnacngncngcngnwsnyntlytygtlncarccmngnayytlngtncntgytnc 1103
Db 121312 TGGGCCAGGCCAGGCTCCCAATGCTGTCTCAGAGCTGAGGACTGTGCTGCATCC 121253

Db	136131	GGAAAGAAATTTCCTAAGCAGCAAAAAACATTCAAGGTGACCTTGGGTCTCTTTAATGATTT	136190
Qy	804	yswnttca/thmgngargcngarcayaarawswnsngaraaytlncaycngayaaytnat	863
Db	136191	CAGTTTTATGAGGAGACACAGCAAGTCAAAAGTTTCAGCAATTCAGCTCAGATGAT	136250
Qy	864	haataaraaayacnttlytsngargnaartlyaaaytngcngcngarathgyethg	923
Db	136251	AGAAAGAAAAAACCCATTTCCTGAGGGAATTCAAACGCTGCACAAATTTGCATAG	136310
Qy	924	yaaygargarytlnaaytlnaayccncargayaaygngaraayaltnsngacntlyga	983
Db	136311	TAACAAAGAGCTGATGTATTATCCGCAAGACAAATGGGGAATAATGTCTCCAGGGCATGTCA	136370
Qy	984	tmgnwsmwncarcarsnaahtaarwsnytngcnttlygmgnccmmgmnaarttgytlyg	1043
Db	136371	GAACTCTTCACAGCAGCCCTCCATCCACAGGCTCGAGAGCCTTGAGAAATGGTTTGG	136430
Qy	1044	ygnacncngnccngnswsnytntlytygtlnarcctcmgmngaytngtncntgyfnc	1103
Db	136431	TGGGGCCAGGCCACAGGTCCTCCATCTGTCTCAGCCTTAGGAGCTTGGTGGCTGCATCC	136490
Qy	1104	ngtlnaaywscngtngcwnsangary-----g	1130
Db	136491	AGCCGCTCTACCACTGGCTGTAAGAGGGCCATGTATAGATCAGCTCAGCTGTGCTTCAG	136550
Qy	1131	ngcwnsnccnaarcenttgcarytlnccwnsngngtngarcngtngngcnaaraarws	1190
Db	136551	AGGGGCGCCCTAAGCCTGGCGAGTTTCCACATGAGTTCAGCTTAGTACACAGAAATC	136610
Qy	1191	nmgnatthgarlytngggarcncncnaltmgnltycaraaraethaygnaaycngtbat	1250
Db	136611	AAGAAATTGAGGCTTGGGAACTTCACCTCGATTTTCAGAAAGTGTATGGAACGCTTGAT	136670
Qy	1251	gccmngnarcaraarttygngtngngtngngtngawswsntngngnacsngcnmgntngt	1310
Db	136671	GCCCAAGCCAGAAAGTTTCTGTGCAGGGCGAGGATCTCATGAGAAACCTCTGCTAGCGACT	136730
Qy	1311	ncaraargnaaytngtngtngggarcncncncaymngntncwnsngngncncwns	1370
Db	136731	GCAGAGAGGAATATGGGGGTGGAAGCCACCATATGAGATGCCCTACTCGGCGACCTTCTAG	136790
Qy	1371	nmngcngtngmgngnswncncncnswsmngnytlnarcarygmgngnwnaachgyws	1430
Db	136791	TGGACCTCTGAGAAAGGGCCATGACCTCCAGACCCCGAATGGTATATCCACTACAG	136850
Qy	1431	nytncarcaygtncncngaraarwsnacsngayancartgyarccngtlnaargncngng	1490
Db	136851	CTTGGACACCTGATGCTCGAAAAAGCTGCAGACACTCAACACAGCCCTGTGAAGCAGCAG	136910
Qy	1491	natggarwsgtlnccntlayaaracngtngtngcngarytlnacnaaracngtngnathta	1550
Db	136911	GAGGAGGGCTGTGCCCTTCACAAAGCCACAGGGGTGAGTGGTCCCAAGACCATGGGACCCA	136970
Qy	1551	ytttynctncaeytgcaygaytngaytngtmgncaygngtlnaargngaycaeytlygngc	1610
Db	136971	CCCTCTGCAATCAGCATGACCTCGAGTGTAGACCTTGACTCAAAAGGAGATCATTTTGGAGG	137030
Qy	1611	nytmgntngtgyaytgcycnaacngnttymgnaactayatzgngcngc-nccnytntlyt	1669
Db	137031	TTTTAAATTTGATTTGCTCTTGGCTGTGGATTTTCGGACTTGCACTGGGCCCTGTAAACCCCTTGT	137090
Qy	1670	tygncarttlytccnttlygnaacngcngtlnatyacncar	1710
Db	137091	TTGGCCAAATTTCTCCATTTTGAACAGCGTATTTTACCAG	137131
RESULT	10		
AC015727	175639 bp	DNA	linear
LOCUS	AC015727/c		
DEFINITION	Homo sapiens chromosome 17 clone RP11-420A6 map 17, WORKING DRAFT		
SEQUENCE	31 unordered pieces.		
ACCESSION	AC015727		

VERSION AC015727.4 GI:10045179
 KEYWORDS HTG: HTGS_PHASE1; HTGS_DRAFT.
 SOURCE human.
 ORGANISM Homo sapiens
 REFERENCE Enayayola, Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 AUTHORS 1 (bases 1 to 175639)
 TITLE Homo sapiens chromosome 17, clone RP11-420A6
 JOURNAL Unpublished
 REFERENCE 2 (bases 1 to 175639)
 AUTHORS Birren,B., Linton,L., Nusbaum,C., Lander,E., Allen,N., Anderson,M., Baldin,J., Barua,N., Beckery,R., Boguslavsky,L., Bouknight,B., Brown,A., Castle,A., Collins,S., Collins,S., Collamore,A., Cooke,P., DeRellano,K., Dewar,K., Domingo,M., Donelan,L., Doyle,M., Ferrelia,P., Fitzhugh,W., Forrest,C., Funke,R., Gage,D., Galagan,J., Gardyna,S., Grant,G., Hagos,B., Heaford,A., Horton,L., Howland,J.C., Johnson,R., Jones,C., Kann,L., Karates,A., Klein,J., Lehoczy,J., Lieu,C., Locke,K., Macdonald,P., Margulis,N., McKean,P., McGurk,J., McKenna,K., McLaughlin,J., Meldrum,J., Morrow,J., Naylor,A., Norman,C.H., O'Connor,T., O'Donnell,P., Peterson,K., Pollara,V., Riley,R., Roy,A., Santos,R., Seery,P., Stange-Thomas,N., Stojanovic,N., Subramanian,A., Talamas,J., Testaye,S., Tirrell,A., Vassiliev,H., Vo,A., Wheeler,J., Wu,X., Wyman,D., Ye,W.J., Zimmer,A. and Zody,M.
 TITLE Direct Submission
 JOURNAL Submitted (17-NOV-1999) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA
 COMMENT On Sep 9, 2000 this sequence version replaced gi:5958075. All repeats were identified using RepeatMasker: Smit, A.F.A. & Green, P. (1996-1997) <http://ftp.genome.washington.edu/RM/RepeatMasker.html>
 ----- Genome Center
 Center: Whitehead Institute/ MIT Center for Genome Research
 Center code: WIBR
 Web site: <http://www-seq.wi.mit.edu>
 Contact: sequence.submissions@genome.wi.mit.edu
 ----- Project Information
 Center project name: L1423
 Center clone name: 420_A-6
 ----- Summary Statistics
 Sequencing vector: M13; M77815; 87% of reads
 Sequencing vector: Plasmid; n/a; 13% of reads
 Chemistry: dye-primer-merman; 4% of reads
 Chemistry: dye-terminator Big Dye; 96% of reads
 Assembly program: Phrap; version 0.960731
 Consensus quality: 161512 bases at least Q40
 Consensus quality: 167866 bases at least Q30
 Consensus quality: 170654 bases at least Q20
 Insert size: 170000; agarose-ep
 Insert size: 172639; sum-of-contigs
 Quality coverage: 4.9 in Q20 bases.
 * NOTE: This is a 'working draft' sequence. It currently consists of 31 contigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as runs of N, but the exact sizes of the gaps are unknown.
 * This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.
 *
 1 13947: contig of 13947 bp in length
 * 13948 14047: gap of 100 bp
 * 14048 14259: contig of 212 bp in length
 * 14260 14359: gap of 100 bp
 * 14360 15039: contig of 680 bp in length
 * 15040 15139: gap of 100 bp
 * 15140 15832: contig of 693 bp in length
 * 15833 15932: gap of 100 bp
 * 15933 16689: contig of 757 bp in length
 * 16690 16789: gap of 100 bp
 * 16790 18092: contig of 1303 bp in length
 * 18093 18192: gap of 100 bp
 * 18193 19993: contig of 1801 bp in length

Db	2689	GAAGCTTCACAGCAGACGCCCTCCCATCATCAGACGGCTGAGGGCTGAGAGAAATGGTTTG	2630
Qy	1044	ygnaengncngcngnwnsnyntlytygtlncarcmmngayytngtncntlygtnc	1103
Db	2629	TGGGCGACAGGCCACGAGGTCCCAATGCTGTCTCCAGCTTAGAGGACTTGGTGCCTGCATCCC	2570
Qy	1104	ngtnaaysnscngtngcnwnsgangr-----g	1130
Db	2569	AGCCGCTTACCATGAGGTGTAAAGGGGCCAATGTAGATAGAGCTCAAGCTGTGGCTTCAG	2510
Qy	1131	ngcnwnscncaarcntlygcarytlncnwsngngtngarcnftngngcnaaraaws	1190
Db	2509	AGGGTGCCTTAAGCTTGCGACGTTCACATAGTCTTAGCTTAGGTATACACAAATGC	2450
Qy	1191	nmgnatlgargtngtggarcncncnaltmgntlycaraaraalthaaygnaayccttgat	1250
Db	2449	AAGATTTGAGGTTTGGGAATCTCCACCTGGATTTTCAGAAATGATGTGTGAAGACGCTTGAT	2390
Qy	1251	gcmmngncaaraartlygcngtlnngtngtngnwsnsltngnnaecnwsngcmngtngt	1310
Db	2389	GCCCGACAGAAATTTGCTGGAGGGGCGAGGATTCATGAGGAAACCTGTGCTAGGGGACT	2330
Qy	1311	ncaraargnaaytngntngtggarcncncaaymgntncnwsngngcncnwsnws	1370
Db	2329	GCACAGAGGAAATGTGGGGTGCAGACCCACCATATGGAATCCCTACTCGGACCTTTTAG	2270
Qy	1371	nmngcngtlnngnmgnwsncncnwsnshsmngyntncaraargnmgnwsncaqaws	1430
Db	2269	TGGAGCTGTGAGAAAGAGGGGCCAATGACCTCCAGACCCAGAAATGTTAGTTCACCTACAG	2210
Qy	1431	nytnarcaaygtlncngaraarawsnacsngayaencarttygcarcngtlnaargcngng	1490
Db	2209	CTTGACACGCTAATCGCTGGAAAAAGCTGCAGACACTCAACACGACGCCGTGAAAGCAGCAG	2150
Qy	1491	natgajarsnftncnctntaaraacngtngtngcnaarytlncnaaraacngtngnatlta	1550
Db	2149	GAGGAGAGGCTGGCCCTCAAAAGCCACAGAGGGGTGAGTTGCCCAAGAGACATGGGAACCA	2090
Qy	1551	yytnytncaaytlycaaygaytlnqaygtlmgncaygngtlnaarmngaycaytlygngc	1610
Db	2089	CCTCTTGACATCAGCATGACCTGAGACTGTGAGACTTGAGATCAAAAGAGATCATTTTGGAGG	2030
Qy	1611	nytmngntlygasytlyccnacsngntlymgnaentlayaltggncngt-ncnynltngyt	1669
Db	2029	TTTAAATTTGATTTGCCCTGTGCTGTGATTTTCGGACTTGCAATGGGCGCTGTAAACCCCTTTGTT	1970
Qy	1670	tygncarttlytycncnttygnaacngcngtltacncaar	1710
Db	1969	TTGGCCAAATTTCTCCATTGTGGAAACGCTGTATTTTAAACCA	1929

```

RESULT 11
AL391823/c
LOCUS      AL391823              181842 bp    DNA        linear      HTG-10-JUL-2001
DEFINITION Homo sapiens chromosome 1 clone RP11-319F11, *** SEQUENCING IN
            PROGRESS ***, 12 unordered pieces.
ACCESSION  AL391823
VERSION    AL391823.9  GI:11995113
KEYWORDS   HTG, HTGS_PHASE1, HTGS_CANCELLED.
SOURCE     human.
            Homo sapiens
REFERENCE  Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
            1 (bases 1 to 181842)
AUTHORS    McIay,K.
TITLE      Direct Submission
JOURNAL    Submitted (09-JUL-2001) Sanger Centre, Hinxton, Cambridgeshire,
            CB10 1SA, UK. E-mail enquiries: humgeny@sanger.ac.uk
COMMENT    requests: clonerequests@sanger.ac.uk
            On Dec 28, 2000 this sequence version replaced gi:11990058.
            ----- Genome Center
            Center: Sanger Centre
            Project code: SC

```

```

Web site: http://www.sanger.ac.uk
Contact: humphreys@sanger.ac.uk
----- Project Information
Center project name: BA15Pfl
----- Summary Statistics
Assembly program: XGAP4; version 4.5
Sequencing vector: M13; M7815; 4% of reads
Sequencing vector: plasmid; 108752; 95% of reads
Chemistry: Dye-terminator ET-amersham; 3% of reads
Chemistry: Dye-terminator Big Dye; 96% of reads
Consensus quality: 174456 bases at least Q40
Consensus quality: 177550 bases at least Q30
Consensus quality: 179392 bases at least Q20
Insert size: 180742; sum-of-coverage
Insert size: 162786; 7.7% error; agarose-fp
Quality coverage: 4.40x in Q20 bases; sum-of-coverage
Quality coverage: 4.89x in Q20 bases; agarose-fp
-----
* NOTE: This is a 'working draft' sequence. It currently
* consists of 12 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
*
1
8897 8996: contig of 8896 bp in length
*
8897 8996: gap of 100 bp
*
8897 22635: contig of 13639 bp in length
*
22636 22735: gap of 100 bp
*
22736 55625: contig of 32890 bp in length
*
55626 55725: gap of 100 bp
*
55726 65440: contig of 9715 bp in length
*
65441 65540: gap of 100 bp
*
65541 88588: contig of 23048 bp in length
*
88589 88688: gap of 100 bp
*
88689 94602: contig of 5914 bp in length
*
94603 94702: gap of 100 bp
*
94703 105410: contig of 10708 bp in length
*
105411 105510: gap of 100 bp
*
105511 107644: contig of 2134 bp in length
*
107645 107744: gap of 100 bp
*
107745 126586: contig of 18842 bp in length
*
126587 126686: gap of 100 bp
*
126687 125610: contig of 28924 bp in length
*
125611 155710: gap of 100 bp
*
155711 173103: contig of 17993 bp in length
*
173104 173203: gap of 100 bp
*
173204 181842: contig of 8639 bp in length.
*
location/Qualifiers
1..101842

```

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misc_feature      1. .8896
                  /note="assembly-fragment:02130
                  clone_end:sp6
                  vector_side:left"
misc_feature      8997. .2265
                  /note="assembly-fragment:01909
                  fragment_chain:1"
misc_feature      22736. .55625
                  /note="assembly-fragment:00961
                  fragment_chain:1"
misc_feature      55726. .65440
                  /note="assembly-fragment:00129
                  fragment_chain:1"
misc_feature      65511. .88588
                  /note="assembly-fragment:00944
                  fragment_chain:1"
misc_feature      88689. .94602
```

```

/note="assembly_fragment:01651
fragment_chain:1"
misc_feature
94703..105410
/note="assembly_fragment:01169
fragment_chain:1"
misc_feature
105511..107644
/note="assembly_fragment:00035
fragment_chain:1"
misc_feature
107745..126586
/note="assembly_fragment:01445
fragment_chain:1"
misc_feature
126687..155610
/note="assembly_fragment:00583
fragment_chain:1"
misc_feature
155711..173103
/note="assembly_fragment:00742
fragment_chain:1"
misc_feature
173204..181842
/note="assembly_fragment:01461"
BASE COUNT 53809 a 39941 c 37882 g 49105 t 1105 others
ORIGIN

Query Match 28.2% Score 488.6 DB 2 Length 181842;
Best Local Similarity 47.5% Pred NO.3.2e-108;
Matches 486; Conservative 212; Mismatches 290; Indels 36; Gaps 2;

```

Query Match	Similarity	28.2%	Score	488.6	DB 2:	Length	181842:
Best Local	Similarity	47.5%	Pred. No.	3.2e+108:			
Matches	486:	Conservative	212:	Mismatches	290:	Indels	36:
						Gaps	2:
QY	743	argargaralhsnaarcrcarcswnathcaargarglnacnsggylunlynlnaargnt	802				
Db	64835	AAGAGGATTTCTTAAGCAGCAAGCAATTTGAGATGTGACTTTGGTGTCTTAAGGCAT	64776				
QY	803	lywsntlyaltnmgargcngarcsyaarawswnsngaraayltncaycncgaayaayltna	862				
Db	64775	TCAGTTTTATAAGGAGCAGATCTATAAAATTTGGAAAAATTTGGACGCTGACTATGCA	64716				
QY	863	lhaaraataaraaycncntlywsngargnaarltlyaaayltngcngcngaraahlygath	922				
Db	64715	TGAGAAAGAAAAACCATTTTCTGGGGAGAAATTCAGACCTTATTCAGAAATTTGCTTA	64656				
QY	923	gyaaaygargarylnaaygltnaaycncnargayaayyngngaraayaaltnwsntgacntlyc	982				
Db	64655	GTAGCAGAGGAGCCTTAATGTTATCCCAAGACCATCGGGGAATGCTCTACAGGCGCTGC	64596				
QY	983	armgwmwmsnccarcarwsnalhaarwsnytnqcnltgmgncnmgng---naarltgt	1039				
Db	64595	AGGGAACCTTCATGCTGCCCTCCCATCCAAAGCTGCAGGCCCAAGGATTAATAAATGCT	64536				
QY	1040	lytgytgnacngnncngnwnsnylnlytgytlncaarcnmngnalytngltnccnlyg	1099				
Db	64535	TTTCATGGGCGCAGGCCAGCGGCTCTGTGGCGGTGTGACAGCAGAGGACTGTGGTCCCTGTG	64476				
QY	1100	tnccng-----tnaaywsngcngltnngcwsng	1126				
Db	64475	TCGCCAGCTGCTCAGACTATGCTGTGAAGGGGCCAAGGTAGAGCTTGGCTGTGGCTTACG	64416				
QY	1127	argyngwmsncnaarcnctlgcaarlynccnwsngngtngarcnngcngngcnaara	1186				
Db	64415	AGGGTCAGAACCCCAAGCCTTGGCAGCTTCCACGTGATTTTAAAGCTGTGGGTGCACAA	64356				
QY	1187	arwsnmgaltnhgarlyntlggarccnccnahtmgntlycaraaralhtlaygnaaycnc	1246				
Db	64355	AGCTCAAGAACTAGGTTTGGGAACCTCTGCTTAAGATTTCAAGAAAGATGTATGAAACGCT	64296				
QY	1247	ggaatccmngncaaraarltlygcngltnngngltnngwmsnswntgngnnaewsnngcmng	1306				
Db	64295	GGATGCCCGGGAAGATTTGCTGTGAAGGTGTGGGACCTTCATGTGAACCTCTGCTAGGG	64236				
QY	1307	tngtlncaraargnaaygltngnltggarccnccncaayngntlnccnwsngngncncmw	1366				
Db	64235	CAGTGCAGAGAAATGTGGGGTCAAGACCCCAACAGATGTCCTACTGGGGCACAC	64176				
QY	1367	swmsnmngcngltnmgngnwsnccnccwnswntgntlyncaraargnmgnwsnagc	1426				

Db	64115	CTAGTGGAGCGTGTAGAAAGGGGGCCACTGTCTCCAGACCCGAAAGTATGATCCACTG	64116
QY	1427	aywsnylnrcarcaayltnccngaraarwsnacsngayacncartltycarccnglnaargcng	1486
Db	64115	ACACGCTTCACACATTCACCTGGAAAGCCACAGACATCAACGGCCGACCTATGAAGACG	64056
QY	1487	cngnratlgtarwslnglnccnlayaaraecnlgntlgtcngarylnaenaaarccnglnnga	1546
Db	64055	CCAGAGGAGAGCGCTTACCCTGCAGAAAGCCACAGGGGTGGAACTGGCCAAAGACTATGGGAA	63996
QY	1547	tlhtaylnlylnaclygcaygaayltnagayltnmgncaygnglnaargngaycaatltyg	1606
Db	63995	CCTACCTCTTGTCATGAGCATGACCTGGATGTGTAGACCTGGTGTCAAAGAGATCATTTTGG	63936
QY	1607	gngcnylnmgntltygayltygcacnacsngnltlmyngacnlayaavgngcnglnccnlynt	1666
Db	63935	AGCTTTAAAAATTTTGACGTGCTCACTGGATTTCAAGACTTTGACGGCCCTGTACACACTTT	63876
QY	1667	gyltlygmcnartltygcntltlygnaecnngcnltltyacncartltyltnlaylncayt	1726
Db	63875	GTTTTGGCCAAATTTCTCCATTTGGAGGGCTATATTATTAACCAATGATTTGAACCCCAATT	63816
QY	1727	g yat 1730	
Db	63815	GTAT 63812	

RESULT	12
AL353634	
LOCUS	77405 bp DNA linear HTG_10-JUL-2001
DEFINITION	Homo sapiens chromosome X clone RP11-435A2 map q21.33-22.3, ***
SEQUENCING	IN PROGRESS.***, In ordered pieces.
ACCESSION	AL353634
VERSION	AL353634.4 GI:9863649
KEYWORDS	HTG; HTGS_PHASE2; HTGS_CANCELLED.
SOURCE	human.
ORGANISM	Homo sapiens
Eukaryota;	Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia;	Eutheria; Primates; Catarrhini; Hominiidae; Homo.
1 (bases 1 to 77405)	
McLay, K.	
Direct Submission	
Submitted (09-JUL-2001)	Sanger Centre, Hinxton, Cambridgeshire,
CB10 1SA, UK. E-mail enquiries:	humquerry@sanger.ac.uk
Requests: clonerequests@sanger.ac.uk	
On Aug 21, 2000 this sequence version replaced gi:9213111.	
COMMENT	

```

Web site: http://www.sanger.ac.uk
Contact: humnery@sanger.ac.uk
-----
Project Information
Center project name: BA435A2
-----
Summary Statistics
Assembly program: XGAP4, version 4.5
Sequencing vector: Plasmid, L08752, 100% of reads
Chemistry: Dye-terminator ET-emersham, 13% of reads
Chemistry: Dye-terminator Big Dye, 86% of reads
Dye-terminator Big Dye: 86% of reads
Consensus quality: 76604 bases at least Q40
Consensus quality: 77098 bases at least Q30
Consensus quality: 77320 bases at least Q20
Insert size: 77405; sum-of-contigs
Insert size: 85298; 1.5% error; agarose-fp
Quality coverage: 4.54x in Q20 bases; sum-of-contigs
Quality coverage: 4.30x in Q20 bases; agarose-fp
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NOTE: This is a 'working draft' sequence.
* This sequence will be replaced
* by the finished sequence as soon as it is available and
* the accession number will be preserved.

Location/Qualifiers
1..77405
/organism="Homo sapiens"

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misc-feature
/ db_xref="taxon:9606"
/ chromosome="xy"
/ map="q21.33-22.3"
/ clone="RP1-435A2"
/ clone_1bp="RPC1-11.2"
1..77405
/note="assembly-fragment:00554"
BASE COUNT      25072 a 14511 c 15133 g 22649 t
ORIGIN
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Query Match	28.2%	Score 488.2	DB 2	Length 77405
Best Local Similarly	47.0%	Pred. No. 3e+108		
Matches 508	Conservative 225	Mismatches 310		Indels 37
				Gaps 4

Oy	668	ytncaycytygaarqthaayalygyltnatgmnaarcathtytlungnaayaaargar	747
Db	8637	ctgcccttgagayctctggaaattttgaaccttgagaaagatgattttaggctatcaggctgaa	8696
Oy	748	garathwsnaaaccarcatwsnathcargargynaactgglytnlytnaargcnclysn	807
Db	8697	gaattttctaaagcaccaaaagcatttcagaaggtgacttgctgctctttaaaggcactcagt	8756
Oy	808	tttathmgargargngarccayaaarwsnswngaraaytlnacymngayaaaytlnahaar	867
Db	8757	ttttgcagaaggaagcagacacattaaaagtttcagaaaaattttcacccctgacacattgtgatgaa	8816
Oy	868	aaaraaycc-nttynswngargnaatttlyaarlungcngngarathtyalttlyaa	926
Db	8817	aaagaaacccctttttcttgatgatpaaattgaagccagctgcagaaattttgcatpaaagtta	8876
Oy	927	ygargarytlnaaygtlnaayccnccargayaayngnaraayaalhsan tgaacttgcarn	986
Db	8877	cgagagacccaattgttaattacctaagacacatgggaaaaatattctocatgcttggtcagag	8936
Oy	987	nwsnswncarcatwsnathhaarwsnytl--cnytgmgncmngmngnaartglttlyg	1044
Db	8937	gctttcacggcgagccccccttcacatctacgtgccccccagagcccttagagagaaaaaggtttat	8996
Oy	1045	gngnagngncngmgwshytltygytlnacrcmngmngaytltungcnytlncn	1104
Db	8997	gggtggggcccgaggtcccatcatgtatgtgcagtctaaaggacttggtgcccgtgtccca	9056
Oy	1105	gtnaa-----ywsngcngtungcwsngargn	1131
Db	9057	gccactccacgcatpactgaamaaggggccaaagttatagcttgggcgcttgcttcaaaagat	9116
Oy	1132	gcwswncnnaarccnhtgcaarytncnwsngngtngarcngtungngcnaaraarwsn	1191
Db	9117	ggaagcccccagcccttggcgagcttcacatgtgttggaagccgcagagcttcacatgaaagtca	9176
Oy	1192	mgnahtgargtbtggarccnccnaltmgnttlyaraarathlyargmaaycncatgatz	1251
Db	9177	aaatattgaagtttgggaacctccacctacatttccagaaagatgtatagaatgccttgagtg	9236
Oy	1252	ccmngnccaraarctlycngtungngtngnswanwsntgmgnaachwsnngcnnngtln	1311
Db	9237	cccagagcaaaatttgctcagagggcgagagacctcatgagaaacctcttttaggggtgg	9296
Oy	1312	caraaragnaaytngntngtggarcnccnccncaymngtlnccwswngngcncnwsnsw	1371
Db	9297	tggaaagggaaaaagtggggtcatagccgcacacagagttcccttaaccagggagccacctagt	9356
Oy	1372	mgngcngtmgmgmwsnccnccnswmngnylncaaraargmgmngmnaacngayln	1431
Db	9357	ggagctgttagaagagggccacacctcttccaaaccccaaaatgtrgatccacccgacac	9416
Oy	1432	ytncaraytlnccngaraaraarwsnccngayaacnartlyrcarcngtlnaargcngcngn	1491
Db	9417	ctgcacacccgtgctccttggaaaagcccgacagacattcatatgccacctcttgaagacagcaggg	9476
Oy	1492	atgcgarwsngtlnccnbaayaaracngtungcngargtlnacnaaraaracngtungnathay	1551

Db	9477	AGGAGGCGCTGTAACCTGCGAAGGCCACAGGGGCGAGAGCTGCCCAAGATCATCTGGGAACCAAT	9536
Qy	1552	ytctyencayctgcaayaytyingaytglmngcayggnglnaarmngaycayletygngcn	1611
Db	9537	CTCTTGATCATGACATGACGTGGATATGAGACATGAGTCACAAAGCAATCATTTTGAGACT	9596
Qy	1612	yltmgnttygatyctgycacnngntlymgnaentayatagggncngt-nccnylnlylt	1670
Db	9597	TTAAGATTTGCATGCTCCACATGAGATCTCAGACTTCACATGGGGCCGTGTGACCTCTTTGTGT	9656
Qy	1671	yggmncatlytlytconcttygnaengcnngtlylancnaryylnhlayencaytgnat	1730
Db	9657	TGGCCCAATATCTCCATTGTGAGATGGCGTATTTATTCACAAAGCTGATACCCCATTTGAT	9716

RESULT	13
HSJ633H17/c	
LOCUS	HSJ633H17
DEFINITION	Human DNA sequence from clone 633H17 on chromosome 1p31.2-32.2.
	124531 bp DNA linear PRI 07-FEB-2000

sequence. Contains a pseudogene similar to part of MRCO1 (cytochrome c oxidase 1), MYCO2 (cytochrome c oxidase II), MTAMP8 (AMP synthase 8) and GOR2 glutamic-oxaloacetic transaminase 2, mitochondrial (aspartate aminotransferase 2) pseudogenes, ESTs and GSSs, complete sequence.

ACCESSION	AL049710
VERSION	AL049710.18
KEYWORDS	GI:5650682 HTG; aspartate aminotransferase; ATP synthase; Cytochrome C oxidase; glutamic-oxaloacetic transaminase; GOT2; MTATP8; MTGOL; MTCO2.
SOURCE	human.

ORGANISM	REFERENCE
Homo sapiens	1 (bases 1 to 124531)
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.	
Moore, M.	

TITLE Direct Submission
JOURNAL Submitted (21-SEP-1999) Sanger Centre, Hinxton, Cambridgeshire, CB10 1SA, UK. E-mail enquiries: humquerry@sanger.ac.uk Clone requests: clonerequests@sanger.ac.uk
COMMENT On Jul 29, 1999 this sequence version replaced g1:556655. During sequence assembly data is compared from overlapping clones

Where differences are found these are annotated as variations together with a note of the overlapping clone name. Note that the variation annotation may not be found in the sequence submission corresponding to the overlapping clone, as we submit sequences with only a small overlap as described above.

This sequence is the entire insert of clone 633H17. This sequence has been finished according to sequence map criteria as follows. An attempt is made to resolve all sequencing problems, such as compressions and repeats, but not necessarily within known annotated human repeat sequence elements (e.g. Alu). Where the

sequence is ambiguous, there is an annotation using the 'unsure' feature key. This sequence was generated from part of bacterial clone contigs of human chromosome 1, constructed by the Sanger Centre Chromosome 1 Mapping Group. Further information can be found at

<http://www.sanger.ac.uk/ncv/cpi1633h17> is from the library RPI-4 constructed at the Roswell Park Cancer Institute by the group of Pieter de Jong. For further details see <http://dnapac.med.buffalo.edu/VECTOR>. pCYPAC2 The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases.

FEATURES
source 1.124531
location/qualifiers
http://www.sanger.ac.uk/Projects/C_elegans/wormpep.
on the WORMPEP database can be found at
EMBL, SWISSPROT, TrEMBL, WPI, WORMPEP. Information
numbers given in the feature table with source database:

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/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="1"
/map="p1.2-32.2"
/clone_lib="RPCI-4"
/clone="633H17"

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repeat_region	1. .234	/note="L2 repeat: matches 2578. .2707 of consensus"
misc_feature	/note="AlusX repeat: matches 59. .292 of consensus"	22083. .22259
repeat_region	/note="match: GSS: Em:AQ055811"	/note="FRAM repeat: matches -2. .175 of consensus"
repeat_region	559. .760	22280. .22393
repeat_region	/note="MIR repeat: matches 74. .262 of consensus"	/note="L2 repeat: matches 2541. .2645 of consensus"
repeat_region	1158. .2018	22394. .22748
repeat_region	/note="LIMA4 repeat: matches 5282. .6145 of consensus"	/note="MLT1B repeat: matches 1. .390 of consensus"
repeat_region	2026. .2590	22749. .22846
repeat_region	/note="HRVL repeat: matches 4676. .5241 of consensus"	/note="L2 repeat: matches 2645. .2745 of consensus"
repeat_region	2879. .9441	22864. .23010
repeat_region	/note="LIPA2 repeat: matches 1. .6146 of consensus"	/note="L2 repeat: matches 2329. .2429 of consensus"
repeat_region	10000. .10124	23011. .23310
repeat_region	/note="MIR repeat: matches 34. .154 of consensus"	/note="Alub repeat: matches 1. .299 of consensus"
repeat_region	10623. .10689	23311. .23391
repeat_region	/note="MIR repeat: matches 36. .102 of consensus"	/note="L2 repeat: matches 2429. .2521 of consensus"
repeat_region	10690. .11014	23433. .23730
repeat_region	/note="L2 repeat: matches 2389. .2710 of consensus"	/note="MST repeat: matches 66. .394 of consensus"
repeat_region	11136. .11262	23731. .25397
repeat_region	/note="L2 repeat: matches 2349. .2473 of consensus"	/note="LIPB1 repeat: matches 4442. .6123 of consensus"
repeat_region	13420. .13522	25477. .25556
repeat_region	/note="MIR repeat: matches 114. .219 of consensus"	/note="MST repeat: matches 1. .85 of consensus"
repeat_region	13602. .13725	25557. .25651
repeat_region	/note="MIR repeat: matches 130. .262 of consensus"	/note="L2 repeat: matches 2585. .2692 of consensus"
repeat_region	13739. .13987	25745. .25785
repeat_region	/note="L2 repeat: matches 2397. .2666 of consensus"	/note="L2 repeat: matches 2702. .2742 of consensus"
repeat_region	14368. .14486	26898. .27373
repeat_region	/note="L2 repeat: matches 2585. .2705 of consensus"	/note="MER31A repeat: matches 1. .661 of consensus"
repeat_region	14656. .14764	27378. .27483
repeat_region	/note="MIR repeat: matches 124. .252 of consensus"	/note="MER93 repeat: matches 292. .397 of consensus"
misc_feature	14765. .14965	27484. .27868
repeat_region	/note="match: GSS: Em:B30406"	27537. .27688
repeat_region	14966. .15150	/note="MER93 repeat: matches 1. .188 of consensus"
repeat_region	/note="MIR repeat: matches 13. .217 of consensus"	27912. .28051
repeat_region	15157. .15818	/note="LIPA13 repeat: matches 6011. .6152 of consensus"
misc_feature	/note="MER39b repeat: matches 14. .579 of consensus"	29900. .30121
repeat_region	16403. .16945	/note="HRVL repeat: matches 3020. .3235 of consensus"
misc_feature	/note="match: GSS: Em:AQ454645"	30122. .30487
repeat_region	16410. .16933	/note="MLT1A2 repeat: matches 1. .374 of consensus"
misc_feature	/note="match: GSS: Em:AQ357205"	30488. .30578
repeat_region	16410. .17037	/note="HRVL repeat: matches 2933. .3020 of consensus"
misc_feature	/note="match: GSS: Em:AQ309712"	30579. .30796
repeat_region	complement(16459. .16861)	/note="MER58A repeat: matches 1. .221 of consensus"
repeat_region	/note="match: GSS: Em:AQ197233"	30797. .31887
repeat_region	16531. .16576	/note="HRVL repeat: matches 1792. .2933 of consensus"
repeat_region	/note="23 copies 2 mer tt 748 conserved"	complement(31888. .32090)
repeat_region	18053. .18154	/note="match: GSS: Em:B58153; match: GSS: Em:B66544;
repeat_region	/note="MIR repeat: matches 7. .107 of consensus"	match: GSS: Em:AQ759038"
repeat_region	18377. .18492	32922. .33453
repeat_region	/note="MER39b repeat: matches 476. .574 of consensus"	32922. .33453
misc_feature	18525. .18954	/note="match: GSS: Em:AQ751315"
repeat_region	/note="match: GSS: Em:AQ210513"	33175. .33412
repeat_region	19026. .19126	/note="L2 repeat: matches 2473. .2708 of consensus"
repeat_region	/note="MIR repeat: matches 10. .114 of consensus"	33454. .33885
repeat_region	19866. .20007	/note="MLT1C repeat: matches 6. .466 of consensus"
repeat_region	/note="MIR repeat: matches 28. .169 of consensus"	33896. .33946
repeat_region	20273. .20334	/note="MIR repeat: matches 162. .215 of consensus"
repeat_region	/note="LTR16C repeat: matches 90. .152 of consensus"	33947. .34302
repeat_region	20403. .20505	/note="THE1B repeat: matches 1. .364 of consensus"
repeat_region	/note="MER81 repeat: matches 1. .110 of consensus"	34303. .34334
repeat_region	20508. .20690	/note="MIR repeat: matches 129. .162 of consensus"
repeat_region	/note="LTR16C repeat: matches 190. .376 of consensus"	34335. .34703
repeat_region	20692. .20725	/note="THE1C repeat: matches 1. .369 of consensus"
repeat_region	/note="L2 repeat: matches 2660. .2693 of consensus"	34704. .34830
repeat_region	20828. .20863	/note="MIR repeat: matches 20. .129 of consensus"
repeat_region	/note="18 copies 2 mer ac 948 conserved"	34831. .35552
repeat_region	21172. .21304	/note="LIMC2 repeat: matches 5573. .6325 of consensus"
repeat_region	/note="MER5A repeat: matches 40. .167 of consensus"	36612. .36657
repeat_region	21201. .21331	/note="L2 repeat: matches 2675. .2723 of consensus"
repeat_region	/note="MER5A repeat: matches 56. .188 of consensus"	36936. .37004
repeat_region	21466. .21528	/note="L2 repeat: matches 2658. .2707 of consensus"
repeat_region	/note="MER54A repeat: matches 214. .285 of consensus"	37383. .37648
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Query Match		
28.2%; Score 488.2; DB 9; Length 124531;		

repeat_region	/note="THE1B element fragment"
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repeat_region	/note="L1 element fragment"
	19720. .19797
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	19877. .19915
repeat_region	/note="MER21B element fragment"
	20007. .20256
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	20518. .20734
repeat_region	/note="MER30 element fragment"
	22661. .22730
repeat_region	/note="MLT1B element fragment"
	22740. .22814
repeat_region	/note="MLT1A element fragment"
	22917. .22974
repeat_region	/note="MLT1A element fragment"
	23342. .23501
repeat_region	/note="THE1B element fragment"
	23550. .23697
repeat_region	/note="THE1B element fragment"
	23725. .24279
repeat_region	/note="THR element fragment"
	24395. .24593
repeat_region	/note="THR element fragment"
	24643. .25270
repeat_region	/note="THR element fragment"
	25277. .25463
repeat_region	/note="THE1B element fragment"
	25528. .25627
repeat_region	/note="THE1B element fragment"
	26081. .26196
repeat_region	/note="MLT1A element fragment"
	26265. .26661
repeat_region	/note="LTR2 element fragment"
	26797. .26898
repeat_region	/note="MSTC element fragment"
	27949. .28241
repeat_region	/partial
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	28319. .28608
repeat_region	/partial
	/note="Alu repeat: matches 308. .1 of consensus"
	30032. .30159
repeat_region	/note="MER5 element fragment"
	31794. .31980
repeat_region	/partial
	/note="Alu repeat: matches 308. .109 of consensus"
	31981. .32073
repeat_region	/partial
	/note="Alu repeat: matches 93. .1 of consensus"
	32232. .32735
repeat_region	/note="MER9 element fragment"
	35168. .35654
repeat_region	/note="MER4B element fragment"
	40640. .40693
repeat_region	/note="27 copies of 2 mer 94 % conserved"
	7879 C 8141 G 13150 T
BASE COUNT	11544 A
ORIGIN	

Query Match	28.28;	Score 487.6;	DB 9;	Length 40714;
Best Local Similarity	47.08;	Pred. NO. 3.3e-108;		
Matches 507; Conservative	225;	Mismatches 311;	Indels 36;	Gaps 4;

Qy	688	yrcaycaytyaaaglynaaaythgylnaagmnaarcaratylngnaayaarar	747
Db	24003	CGCCCCAAGATCTGTGGATTACCTTGAGAGATGATTATGGGTATCAGGTGAA	24062
Qy	748	garathwnaaccarcacarnmaahcargatynaactggtntynynaargcnlttysn	807
Db	24063	GAATTTCTTAACCAACCAAGCTTCTAAGAGGATCTTGTCCTTTAAAGCACTTCAGT	24122

[illegible]

RESULT	15	
AP004219/c		
LOCUS	159712 bp	DNA
DEFINITION	Hom sapiens genomic DNA, chromosome 8q33, clone: R81223D11,	linear
complete sequence.		
ACCESSION	AP004219	
VERSION	AP004219.2	GT:18146615
KEYWORDS		

SOURCE	ORGANISM	REFERENCE	AUTHORS	TITLE	JOURNAL	COMMENT	FEATURES
repeat_region	Homo sapiens pre-pro-B cell cell_line:FLB 14 - 14 DNA, clone_lib:Keio BAC library clone:KBI222D11.	1 (bases 1 to 159712)	Shimizu, N. and Asakawa, S.	Submitted (28-SEP-2001) Nobuyoshi Shimizu, Keio University, School of Medicine, Molecular Biology, 35 Shinanomachi, Shinjuku-ku, Tokyo 160-8582, Japan (E-mail:nshimizu@med.keio.ac.jp, Tel:81-3-3351-2370, Fax:81-3-3351-2370)	On Jan 14, 2002 this sequence version replaced gi:15824050.	/evidence-not_experimental	12507..12560
repeat_region	Homo sapiens	1 (bases 1 to 159712)	Shimizu, N. and Asakawa, S.	Submitted (28-SEP-2001) Nobuyoshi Shimizu, Keio University, School of Medicine, Molecular Biology, 35 Shinanomachi, Shinjuku-ku, Tokyo 160-8582, Japan (E-mail:nshimizu@med.keio.ac.jp, Tel:81-3-3351-2370, Fax:81-3-3351-2370)	On Jan 14, 2002 this sequence version replaced gi:15824050.	/rpt_family="TAAAAA)n"	/rpt_family="TAAAAA)n"
repeat_region	Mammalia: Eutheria: Primates: Catarrhini: Homnidae: Homo.	1 (bases 1 to 159712)	Shimizu, N. and Asakawa, S.	Submitted (28-SEP-2001) Nobuyoshi Shimizu, Keio University, School of Medicine, Molecular Biology, 35 Shinanomachi, Shinjuku-ku, Tokyo 160-8582, Japan (E-mail:nshimizu@med.keio.ac.jp, Tel:81-3-3351-2370, Fax:81-3-3351-2370)	On Jan 14, 2002 this sequence version replaced gi:15824050.	/evidence-not_experimental	12793..12825
repeat_region						/rpt_family="AT_rich"	/rpt_family="AT_rich"
repeat_region						/evidence-not_experimental	15792..15852
repeat_region						/rpt_family="AluU/FRAM"	/rpt_family="AluU/FRAM"
repeat_region						/evidence-not_experimental	15993..16013
repeat_region						/rpt_family="AT_rich"	/rpt_family="AT_rich"
repeat_region						/evidence-not_experimental	16014..16312
repeat_region						/rpt_family="AluSg"	/rpt_family="AluSg"
repeat_region						/evidence-not_experimental	17126..17436
repeat_region						/rpt_family="AluSg1"	/rpt_family="AluSg1"
repeat_region						/evidence-not_experimental	17798..17960
repeat_region						/rpt_family="L2"	/rpt_family="L2"
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repeat_region						/evidence-not_experimental	19346..19541
repeat_region						/rpt_family="L2"	/rpt_family="L2"
repeat_region						/evidence-not_experimental	19687..19844
repeat_region						/rpt_family="L1MBC"	/rpt_family="L1MBC"
repeat_region						/evidence-not_experimental	19854..20081
repeat_region						/rpt_family="L2"	/rpt_family="L2"
repeat_region						/evidence-not_experimental	20082..20106
repeat_region						/rpt_family="AT_rich"	/rpt_family="AT_rich"
repeat_region						/evidence-not_experimental	20423..20566
repeat_region						/rpt_family="FRAM"	/rpt_family="FRAM"
repeat_region						/evidence-not_experimental	20567..20636
repeat_region						/rpt_family="GA-rich"	/rpt_family="GA-rich"
repeat_region						/evidence-not_experimental	20973..21282
repeat_region						/rpt_family="AluV"	/rpt_family="AluV"
repeat_region						/evidence-not_experimental	22379..22724
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repeat_region						/evidence-not_experimental	22933..23019
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repeat_region						/evidence-not_experimental	23044..23270
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repeat_region						/evidence-not_experimental	23275..23586
repeat_region						/rpt_family="AluSg1"	/rpt_family="AluSg1"
repeat_region						/evidence-not_experimental	23946..24035
repeat_region						/rpt_family="MER94"	/rpt_family="MER94"
repeat_region						/evidence-not_experimental	

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OM nucleic - nucleic search, using sw model

Run on: July 8, 2002, 16:20:51 ; Search time 481.64 Seconds
(without alignments)
6170.340 Million cell updates/sec

Title: US-09-997-610-7

Perfect score: 1731

Sequence: 1 atgynachargantaygg.....gytntatytncaygtatg 1731

Scoring table:

IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 1736436 seqs, 858457221 residues

Total number of hits satisfying chosen parameters: 3472872

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

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24: /SID55/gcgdata/geneseq/geneseqn-emb1/NA2002.DAT:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	471.4	27.2	2590	22	AAH14327
2	452	26.1	3093	22	AAH14327
3	452	26.1	9236	22	AAH14327
4	452	26.1	9236	22	AAH14327
5	452	26.1	9236	22	AAH14327
6	429.6	24.8	3203	22	AAH14327
7	427.2	24.7	6063	22	AAH14327
8	427.2	24.7	31529	22	AAH14327

10	426.4	24.6	6063	22	AAH14327	Human reproductive
11	421.4	24.3	1736	22	AAH14327	Human CDNA sequenc
12	421	24.3	465237	24	AAH14327	Human oestrogen re
13	409	23.6	1485	23	AAH14327	DNA encoding novel
14	408.8	23.6	1278	22	AAH14327	Human secreted pro
15	400.4	23.1	1580	22	AAH14327	DNA encoding novel
16	395.2	22.8	7726	22	AAH14327	Human nervous syst
17	393.6	22.7	32127	22	AAH14327	Human excretory re
18	393.6	22.7	32127	22	AAH14327	Human kidney relat
19	393.6	22.7	72215	22	AAH14327	Human immune/haema
20	388.2	22.4	1405	22	AAH14327	Human immune/haema
21	386.6	22.3	1405	22	AAH14327	Human nervous syst
22	386.4	22.3	8387	22	AAH14327	Human immune/haema
23	386.4	22.3	8387	22	AAH14327	Human reproductive
24	384.4	22.2	1621	23	AAH14327	DNA encoding novel
25	378.8	21.9	3049	22	AAH14327	Human CDNA sequenc
26	373.8	21.6	81369	21	AAH14327	Human T gene DNA.
27	371.6	21.5	1306	23	AAH14327	DNA encoding novel
28	361.6	20.9	22756	22	AAH14327	DNA encoding human
29	361.6	20.9	22756	22	AAH14327	Human reproductive
30	355.4	20.5	1946	22	AAH14327	Human breast cell
31	355.4	20.5	1946	22	AAH14327	Human foetal liver
32	355.4	20.5	1946	22	AAH14327	Probe #2925 for ge
33	355.4	20.5	1946	22	AAH14327	Human brain expres
34	355.4	20.5	1946	22	AAH14327	Human bone marrow
35	355.4	20.5	1946	22	AAH14327	Probe #2908 for ge
36	355.4	20.5	1946	22	AAH14327	Probe #2884 used t
37	355.4	20.5	1946	22	AAH14327	Human immune/haema
38	347	20.0	26410	22	AAH14327	Human immune/haema
39	346.8	20.0	32986	22	AAH14327	Human immune/haema
40	346.8	20.0	32986	22	AAH14327	Human immune/haema
41	346.8	20.0	923	23	AAH14327	DNA encoding novel
42	341.4	19.7	50000	24	AAH14327	Human glutamate re
43	341.4	19.7	50000	24	AAH14327	Human GMS gene tr
44	337.4	19.5	57728	22	AAH14327	Human 9p11 chromos
45	336.6	19.4	2197	22	AAH14327	Human CDNA sequenc

ALIGNMENTS

RESULT 1	
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ID	AAH14327 standard; CDNA: 2590 BP.
AC	AAH14327:
DT	26-JUN-2001 (first entry)
XX	
DE	Human CDNA sequence SEQ ID NO:11697.
XX	
KW	Human: primer: detection; diagnosis; antisense therapy; gene therapy; ss.
OS	Homo sapiens.
XX	
PN	EP1074617-A2.
XX	
PD	07-FEB-2001.
XX	
PF	28-JUL-2000; 2000EP-0116126.
XX	
PR	29-JUL-1999; 99JP-0248036.
PR	27-AUG-1999; 99JP-0300253.
PR	11-JAN-2000; 2000JP-0118776.
PR	02-MAY-2000; 2000JP-0183767.
PR	09-JUN-2000; 2000JP-0241899.
PA	(HELI-) HELIX RES INST.
XX	
PI	Ota T, Isogai T, Nishikawa T, Hayashi K, Saito K, Yamamoto J;
PI	Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;
XX	
DR	WPI; 2001-318749/34.

```
XX Primer sets for synthesizing polynucleotides, particularly the 5602
PT full-length cDNAs defined in the specification, and for the detection
PT and/or diagnosis of the abnormality of the proteins encoded by the
PT full-length cDNAs -
XX
XX Claim 8: SEQ ID 11697, 2537pp + CD ROM; English.
XX
CC The present invention describes primer sets for synthesizing 5602
CC full-length cDNAs defined in the specification. Where a primer set
CC comprises: (a) an oligo-dT primer and an oligonucleotide complementary
CC to the complementary strand of a polynucleotide which comprises one of
CC the 5602 nucleotide sequences defined in the specification, where the
CC oligonucleotide comprises at least 15 nucleotides; or (b) a combination
CC of an oligonucleotide comprising a sequence complementary to the
CC complementary strand of a polynucleotide which comprises a 5'-end
CC sequence and an oligonucleotide comprising a sequence complementary to a
CC polynucleotide which comprises a 3'-end sequence, where the
CC oligonucleotide comprises at least 15 nucleotides and the combination of
CC the 5'-end sequence/3'-end sequence is selected from those defined in
CC the specification. The primer sets can be used in antisense therapy and
CC in gene therapy. The primers are useful for synthesizing polynucleotides,
CC particularly full-length cDNAs. The primers are also useful for the
CC detection and/or diagnosis of the abnormality of the proteins encoded by
CC the full-length cDNAs. The primers allow obtaining of the full-length
CC cDNAs easily without any specialised methods. AAH03166 to AAH13628 and
CC AAH13633 to AAH18742 represent human cDNA sequences; AA892446 to
CC AA895893 represent human amino acid sequences; and AAH13629 to AAH13632
CC represent oligonucleotides, all of which are used in the exemplification
CC of the present invention.
XX
SQ Sequence 2590 BP; 722 A; 526 C; 670 G; 672 T; 0 other;
XX
Query Match 27.2%; Score 471.4; DB 22; Length 2590;
Best Local Similarity 47.0%; Pred. No. 2e-110;
Matches 481; Conservative 213; Mismatches 293; Indels 37; Gaps 3;
XX
QY 744 rgargarathwsnaarcarcarwsnathcarargatgnaentggtntynlnaargntt 803
DB :|||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
DB :|||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
QY 804 ywsnttlyatmgngargngargcayaaarwsnsgaraaytlncaycngsngayaaygnat 863
DB :|||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
DB 668 cagtttcaaaaggaagaaagacataaaatcagaataattgtacgctgtacatgaat 727
QY 864 haaraaraaraaycnttlywsngargnaartlyaaarlyngcngcngarathlythlyb 923
DB :|||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
DB 728 agaaaaagaaaaaacagtttctgtgagagaatctcaagcagctgcaagaatctgtataag 787
QY 924 yaaygaargaytlnaaygtncarcargayaayggngarayaathwsnttgactgyca 983
DB :|||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
DB 788 catcaagaagcctaattgtlaaccacagacatlgggaaatgtctcagcgccatgca 847
QY 984 rmngwsnswnaarcarwsnatharwsnlytngcntgmgngcnmngmng---naatgggt 1040
DB :|||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
DB 848 gagaccttcaacagagccctctcgtcacaagccagagacatccagagaaaaaatgggt 907
QY 1041 ytyygnacngcngcngnswntlytgytncarcnmngnayaaytngtncnltgygt 1100
DB :|||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
DB 908 ttatggccagcagcaggtccctgtgtgtgagcactatagacttggtccctgtgt 967
QY 1101 nccngtlnaayws-----ngcngtngcswnga 1127
DB :|||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
DB 968 cccagctgtctcagcagcatggtcgtgaaagggccaatgtagagctcatgctgtgtctcaga 1027
QY 1128 rggngcwnsncnaarcntgtgcarytncmwsngngtngcngtngcngngcnaaraa 1187
DB :|||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
DB 1028 ggggtgaagccccaagcctgtgcacatccacatggtgtgtgagctgtatagtaaccaaga 1087
QY 1188 twsnmgatlgargtntgggarcncnatlmngntlycaraaraathtlaygnaaycngt 1247
DB :|||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
DB 1088 gtcagaatttggtgttggaacctccacactatgatttcagaagaatgtatggaatgacctg 1147
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QY 1248 gatccmngncaraarhtlycngtngngtngngnswntlgmngnacwnsngcnmngt 1307
DB :|||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
DB 1148 gatgccaggaagaaagtcttgcagtgcaagggccctcatgtgaaacctctgtgagggc 1207
QY 1308 ngtncaraargnaaygtngtngtngarcncncncaymngntncwnsngnccnccns 1367
DB :|||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
DB 1208 aatgcagaaggaattgtgtgggcccagagctccaccagtagagctccctaattggtgcatatc 1267
QY 1368 nwsrmngcngtngmngmngnswncncnswnsnmngntncaraarargmngnswnaanga 1427
DB :|||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
DB 1268 tagtggagctgtggaagaggggccaccgctcttcaagaccacgaatgggtgatatccatga 1327
QY 1428 ywsnlytncarcaytlnccngaraarwsnawcnayacncartgyarcngtlnaargcngc 1487
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DB 1328 cagcttgcacatgcgctgtgaaagacgcagacatccatgccaagctgtgaagaagc 1387
QY 1488 nggnatggarwsngtncntlayaaraengtngtngcngarytlnaaraengtngnat 1547
DB :|||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
DB 1388 caggagggaggtctgtacccttcaagccagcagggcagaactgcgaagcatalggtlcc 1447
QY 1548 htaytlytncaytgycaaygylngaygtmngncaygngtlnaarmngnayaaytlygg 1607
DB :|||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
DB 1448 ccaccttgcatalcagcgtgacccgtaatgtlgagacctagatcaagaagatcatlttgg 1507
QY 1608 ngcnytmngntlygaytgcacnaengntlymgnaentlayatggngcng-tncnylnt 1666
DB :|||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
DB 1508 agcttcaaatgtatgtcctgtgacttgcattgtgaactgtcatgctgtgagccctt 1567
QY 1667 gtytlygcncarttlycncnttlygnacngcngtntlyacncarttlytlnaytlnayt 1726
DB :|||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
DB 1568 gtttggccaatgtctccatcttggaaatgctgtatctaccacatccatgaccccatc 1627
QY 1727 gyat 1730
DB :|||
DB 1628 gtat 1631
XX
RESULT 2
AA503687/c
ID AA503687 standard; DNA; 3093 BP.
XX
XX AA503687;
AC
XX
XX 29-AUG-2001 (first entry)
DE
XX
XX Rhesus gene locus: RHD gene deletion in Rh negative haplotypes.
DE
XX Rhesus box: RHD positive; sequence length polymorphism; SSP; RHD; SMP1;
KW RHE: Rh negative; blood group typing; blood transfusion; antigen C;
KW haemolytic disease of the newborn; chromosome 1 p34.1-p36; ds.
OS
XX Homo sapiens.
XX
XX Key Location/Qualifiers
FH primer_bind 32..54
FT :tag= a
FT /note= "Binding site of primer rex7"
FT primer_bind complement (3034..3054)
FT :tag= b
FT /note= "Binding site of primer rmb31"
XX
XX WO200132702-A2.
XX
XX 10-MAY-2001.
XX
XX 31-OCT-2000; 2000WO-EP10745.
XX
XX 02-NOV-1999; 99EP-0121686.
XX
XX 31-MAY-2000; 2000EP-0111696.
XX
XX (DRKB-) DRK BLUTSPENDEDIENST BADEN WUERTEMBERG.
```


PI	Fliegel WA, Wagner FF.
DR	WPI; 2001-291052/30.
XX	
PT	New nucleic acid molecular structure, useful for detection of common
PT	RHD positive haplotypes in D-negative individuals, comprises RHD, SMPL
PT	and RHCE genes -
XX	
PS	Example 10; Fig 5; 135bp; English.
XX	
CC	The sequence represents the coding sequence of Rhesus gene locus:
CC	RHD gene deletion in Rh negative haplotypes. The Rhesus genes
CC	locus comprises the RHD, SMPL and RHCE (all undefined) genes and/or the
CC	Rhesus box(ss), preferably the hybrid Rhesus box, the upstream Rhesus box
CC	and/or the downstream Rhesus box. The RHD and RHCE genes are located at
CC	chromosome 1 p24.1-p36. Rhesus box flanks the breakpoint region of the
CC	RHD deletion in the common RHD negative haplotypes. The sequence has
CC	been used to design primers which are useful for: (1) the specific
CC	detection of the common RHD positive haplotypes in D-negative
CC	individuals; (2) blood group typing; (3) determining whether a patient
CC	can be transfused with RHD negative blood and whether blood is suitable
CC	for transfusion to patients who should not be exposed to antigen C; (4)
CC	assessing the risk of a Rhd negative mother of conceiving or carrying an
CC	RHD positive foetus. Anti-D antibodies are useful for treating pregnant
CC	women who are Rhesus D negative, where the foetus is not homozygous for
CC	the RHD gene to treat or prevent haemolytic disease of the newborn.
XX	
XX	Sequence 3093 BP; 891 A; 754 C; 619 G; 829 T; 0 other;

Query Match	26.1%	Score 453;	DB 22;	Length 3093;
Best Local Similarly	47.4%;	Pred. No. 2.4e-105;		
Matches 485; Conservative	212;	Mismatches 282;	Indels 45;	Gaps 5;

Qy	744	rgqraethwmsnrcarcarwmsntheaargtnacntbggfnynlynhaegent	803
Db	1338	AGGGAAATTTCTTAAGCAGCAAAAGCATTCACAGAGGAGACTGGGGTGTCTTAAAGACATT	1279
Qy	804	ywnstlyethlmngargarvcngarceayaarwsnmsngaraaylencycngsyaagtnat	863
Db	1278	CAGTTTATTAAGGGAAGCAGACATTAAGTTCCGGAAATTTCCAGCCTGCACATTTGAT	1219
Qy	864	haaraaraaraa--yccntltwysngargnaarlttyaarylngcngcnagaraathlyath	921
Db	1218	AGAAAGAAATAATTCATTTCTGTGAGGAGAAATTCAGACTGGCTGCAGAAATTTGCATG	1159
Qy	922	tyyaaygararyltynaayltnaaylncncnargyaaygngnagaraayathwstlbgacntly	981
Db	1158	AGTAAC-AGGAGCAATATGCTAATTTCCCAAGACATGGGAAATGTCCTCAGGGCATGT	1100
Qy	982	carngnwsnscarcaryaethaawaanyng-ctngbmncnmngmaarctgtc	1040
Db	1099	CAGAGCTTTATTAAGGCACACCCCTCCCATCAGAGTTCAGAGTATCAAGAAAAATGTT	1040
Qy	1041	yltygnaecnngnccngnawsnylntlytlygltnaarcnmngnalyentlncnltgvt	1100
Db	1039	TTGTTGGCCAGGCCGGGGCTCATGCTGTGACACCTAGGACTTGGTGCCCTGAT	980
Qy	1101	ncngntna-----aywsngcngltngcwsng	1126
Db	979	CCCGACACCTCCCAACCATGACTGACGGGAGGCAAGCATAGACCTTGGCGCTTACTTCGCG	920
Qy	1127	argngcwnscnnaarccntlggarctyltncnwsngngltngarcngtngngcnara	1186
Db	919	GGATGTCAAGCCCAACCTTGACACCTTCCATGTGGTGTTCAGACTCGAGTGCACAGA	860
Qy	1187	arwsmgnaethgarlntlggarccnccnalthmgnltlycaraaazlthcaygnaaycmt	1246
Db	859	AGTCAAGAAATTTGGGGTTTGGAACCTTCCCTAAGATTAAAGAGATGTGGCGAATGCT	800
Qy	1247	ggaatgcnmngcnaaraatlygcngltngngltngnwsnslntgmgnaenwsngcmngng	1306
Db	799	GGATGGCCCATCGAAGATTGTCTGCAGGGAGCGGGCCCTCATAGAGATCTCTCTCCAGGG	740

Qy	1307	ttgtctcacaagaagaaatgtgtgntgttgagatgccnccncaatgmgnttncnswngngcncw	1366
Dy	739	CAGTGCAGAAAGGAAATGTGGGGTCAGAGACCCACACACAGTCCCTACTGTGGGACAC	680
Qy	1367	swswtmngcgtctumgmgnwscnccnswtmngnylncaaraagmgnwsnacng	1426
Dy	679	CTAGTGGAGCGTGTAGAAAGAGTGTCTCC-----AGACCCAGATGGTATAGTCCACGG	627
Qy	1427	aywsnylncaatcaygtlncngaraarwsnacngayacncaatgtgcacnglnaargcng	1486
Dy	626	ACAGCTTGACACCGTGTACCTCGGAAAACCTCAACACTCAATGTCCAGCCATGAAGAG	567
Qy	1487	cngnnaatgawsgtnccttayaaracngtngtngcngaylnaacaaracngtngna	1546
Dy	566	CTGAGAGGGAGCGTGTACCTCGCAAAAGCTACAGGGCAGACAGCTGCCCAAGACATGGGA	507
Qy	1547	tlhtayltngcayctgyccaygaytngaygtlmgncaygnglnaarimgncaatctgy	1606
Dy	506	CCCAACCCCTTCATCTCGTGGACCTGGATGTGATGTGAGTCAAGAGAGATCATTTTG	447
Qy	1607	gngcnltmngtltgaytlygcnaacngntltmgnaentlaytbgncnglncnynt	1666
Dy	446	GAGCTTTAAGTTTGAATGTGCCCACTGGATTTTGGACTCTCATGGGCGTGTAGCTCTTT	387
Qy	1667	gyltlygncatctlycnclytlygnaacngcnclytaacncaatlyylnatylncat	1726
Dy	386	GTTTGAACCATTTATCCATTTGGAAATGGCTGTATTTACCCAAATGCCGTATGCCCAATT	327
Qy	1727	gyatc 1730	
Dy	326	GTAAT 323	

RESULT	3
AA503689/c	
ID	AA503689 standard; DNA: 9236 BP.
XX	
AC	AA503689;
XX	
DT	29-AUG-2001 (first entry)
XX	
DE	Rhesus gene locus: upstream Rhesus box of D-positives.
XX	
KM	Rhesus box: RHD positive; sequence length polymorphism; SSP; RHD; SMP1.
KM	RHC: Rh negative; blood group typing; blood transfusion; antigen C;
KM	haemolytic disease of the newborn; chromosome 1 p34.1-p36; ds.
XX	
OS	Homo sapiens.
XX	
PN	W0200132702-AZ.
XX	
PD	10-MAY-2001.
XX	
PF	31-OCT-2000; 2000WO-EP10745.
XX	
PR	02-NOV-1999; 99EP-0121686.
PR	31-MAY-2000; 2000EP-0111696.
XX	
PA	(DRKB-) DRK BLUTSPENDEDIENST BADEN WUERTTEMBERG.
XX	
PI	Flegel WA, Wagner FF;
XX	
DR	WPI: 2001-291052/30.
XX	
PT	New nucleic acid molecular structure, useful for detection of common
PT	RHD positive haplotypes in D-negative individuals, comprises RHD, SMP1
PT	and RHC genes -
XX	
PS	Disclosure: Fig 9; 135pp; English.
XX	
CC	The sequence represents the coding sequence of Rhesus gene locus: upstream Rhesus box of D positives. The Rhesus genes locus

D	b	6014	AGGGAATAATTCTTAACAGCAGCAAGCACTTCAAGAGGTGACTTGGGTGTGCTTTAAAGACATT	59535
Q	y	804	ywnshlyatlmgngarqcngarcaayaarwswnsngaraaylylncayccngayaayltnat	863
D	b	5954	CAGTTTATATAAGGAAAGCAGAGACTAAAGATTCCGAAATTTGGACCTGCACAAATGTAT	58955
Q	y	864	haaraataaraa--ycentlywsngaragmnaartlyaaarylthngcngcngrataltgath	921
D	b	5694	AGAAATAAATAATTTCCATTCTTTCGTGAGGAGAAATTCACACTGGCGCAAAATTTGCA	58353
Q	y	922	tyayaaygarjayltnaaygtlnaayccncaragyaaygaragrarataltwstltnatgacny	981
D	b	5834	AGTAAAC-AGGAGCCAAATGTCTTAFTTCCCAAGACAAATGGGAAATGTCTCCAGGGCATGT	57766
Q	y	982	carngnswnsncarcarwshthaarwanlyng-ctlygmngcnmgmgnaartgylt	1040
D	b	5775	CAGAGGCTTTATAGGCCAACCCCTCCCATCAGAGTCCAGAGGTATCAGGAAATAATG	57186
Q	y	1041	yltyygnacngnmgcnngnswsnylntygtfgytlnearccnmngsyylnthncntlygt	11000
D	b	5715	TTTGTTCACAGGGCCGGGGTTCATGCTGTGTGACCTAGGAGACTTGTGTCCCTGCAT	56566
Q	y	1101	ncngttha-----aywsnngtngcnwng	11266
D	b	5655	CCGAGCCACTTCCACCATCTACTGACGGGAGGCAAGTACAGACTTGGGCTGTAGCTTGG	55966
Q	y	1127	argyngcnwsncnaarcnltggcarylncnwsngnglngarcnglngngcnaara	11866
D	b	5595	GGAGTTCAGAGCCCAACCTTGCACGCTTCCATGTGTGTGACTGGAGTGCAGATGCACAGA	55386
Q	y	1187	arwsnmnatltgatrltltggarccnccnatlmngtlyaratarathtaygnaaycent	12466
D	b	5535	AGTCAAAATTTGGGTTTGGAAACCTTCGCTAGATTAAGAGGATGTGCGGAAATGTCT	54766
Q	y	1247	ggatcgcnmgncaraarctlygcngtngnglngngswnsntlygmngacnswngnmng	13066
D	b	5475	GGATGCCAGTCCAGAGATTGTCTGTCAGAGAGAGGGCCCTCATGTGAGATCTTCTGCCAAG	54186
Q	y	1307	ltnlncaraaragnaayltnnglntggarccnccncaymnglncnwsngngcncn	13666
D	b	5415	CAGTGCAGAAAGGAAATGTGGGGTCAAGAGACCCACACACAGTCCCTACTGGGGCACAC	53586
Q	y	1367	snwsnmngcnngltmgmgmngswnsncnccnswnsnmngylncaraaragmgmngnacng	14266
D	b	5355	CTAGTGGAGCTGTGAGGAAGAGTCTCTC-----AGACCCACAGATGTGATGATCCACCG	53036
Q	y	1427	aywsnlnncarccaylthccngaraarwsnccngayacnartgyccatgcnngfnaarng	14866
D	b	5302	ACAGGTTGCACCGTGTACTGTGGAAACCTGCACACACTCAATGCCAGCCCATGAAGACG	52436
Q	y	1487	cngnatltyagwsnnglthccntlayaaraenglnglncngaryltnaenaarcnngtngna	15466
D	b	5242	CTGAGAGSGAGAGCTGTACCTGCACAAAGTACAGGGGCAAGGCTGCCCAAGACATGGGA	51886
Q	y	1547	thtalytynlncaylcyrcaygaylytngayltnmgncaygngltnaarmngsacaytlyg	16066
D	b	5182	CCCAACCCCTTCATCTCGTACGACTGTGATGTGAGATGTGAGTCAAGAGACATCAATTTTG	51226
Q	y	1607	ngcnlytmngltlygarytlycncacngngltnymgnaentlayatggncnglncnlynt	16666
D	b	5122	GAGCTTAAAGATTGTACTGTGCCCACTGTGATTTTGGACTCTCAAGGGCGCTGTAGCCCTTT	50666
Q	y	1667	gytlycngacatlytlycncntlyygnacngnltntlyencarltlyntlaylncayt	17226
D	b	5062	GTTTGGACCAATTATCCCATTTGGAAATGCGCTATATTTACCAATGCCGTACCCCATTT	50036
Q	y	1727	gyat l730	
D	b	5002	GTAT 4999	
RESULT		6		
AAAD05134				

ID	AD05134	standard; cDNA; 3203 BP.
AC	AD05134:	
XX		
DT	17-JUL-2001	(first entry)
DE	Human secreted protein-encoding gene 14 cDNA clone H1SBF60, SEQ ID NO:24	
XX		
KW	Human; secreted protein; proliferative disorder; cancer; tumour;	
KW	foetal abnormality; developmental abnormality; haematopoietic disorder;	
KW	immune system disorder; AIDS; autoimmune disease; rheumatoid arthritis;	
KW	inflammation; allergy; neurological disorder; Alzheimer's disease;	
KW	Parkinson's disease; cognitive disorder; schizophrenia; asthma;	
KW	skin disorder; psoriasis; sepsis; diabetes; atherosclerosis;	
KW	cardiovascular disorder; angiotensin disorder; kidney disorder;	
KW	gastrointestinal disorder; pregnancy-related disorder;	
KW	endocrine disorder; infection; wound healing; vulnerability;	
KW	cell culture; chemotaxis; food additive; gene therapy;	
XX	binding partner identification; ss.	
OS	Homo sapiens.	
XX		
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FH	CDS	153..356
FT		/*tag= a
FT		/product= "Human secreted protein"
FT	sig_peptide	153..215
FT		/*tag= b
FT	mat_peptide	216..353
FT		/*tag= c
FT		/product= "Human mature secreted protein"
XX		
PN	WO200134769-A2.	
XX		
PD	17-MAY-2001.	
XX		
PR	01-NOV-2000; 2000WO-US30040.	
XX		
XX	05-NOV-1999; 99US-0163580.	
PR	30-JUN-2000; 2000US-0215130.	
XX		
PA	(HUMA-) HUMAN GENOME SCI INC.	
XX		
PI	Ruben SM, Komatsoulis GA, Wei P, Baker KP, Fiscella M,	
XX		
DR	WPI; 2001-308781/32.	
DR	P-PsDB; AA01245.	
PT	New isolated nucleic acid molecule encoding a human secreted protein 1s	
XX	used in preventing, treating or ameliorating a medical condition -	
PS	Claim 1; Page 408-409; 519pp; English.	
XX		
CC	AAD05121-AAD05203 represent cDNAs corresponding to 24 human secreted	
CC	protein genes, and AAE0132-AAE01311 represent the proteins they encode.	
CC	AAE01312-AAE01340 represent human secreted protein variants or fragments	
CC	The secreted proteins and their genes are useful for preventing,	
CC	treating or ameliorating medical conditions, e.g., by protein or gene	
CC	therapy. Pathological conditions can be diagnosed by determining the	
CC	amount of the new protein in a sample or by determining the presence of	
CC	mutations in the new genes. Specific uses are described for each of the	
CC	24 genes, based on the tissues in which they are most highly expressed,	
CC	and include developing products for the diagnosis or treatment of	
CC	proliferative disorders, cancer, tumours, foetal and developmental	
CC	abnormalities, haematopoietic disorders, diseases of the immune system,	
CC	AIDS, autoimmune diseases (e.g., rheumatoid arthritis), inflammation,	
CC	allergies, neurological disorders (e.g., Alzheimer's disease,	
CC	Parkinson's disease), cognitive disorders, schizophrenia, asthma,	
CC	skin disorders (e.g., psoriasis), sepsis, diabetes, atherosclerosis,	
CC	cardiovascular disorders, angiotensin disorders, kidney disorders,	
CC	gastrointestinal disorders, pregnancy-related disorders, endocrine	
CC	disorders, and infections. The proteins can also be used to aid wound	
CC	healing and epithelial cell proliferation, to prevent skin aging due to	


```
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PR 01-SEP-2000; 2000US-0229345.
PR 05-SEP-2000; 2000US-0229509.
PR 05-SEP-2000; 2000US-0229513.
PR 06-SEP-2000; 2000US-0230437.
PR 06-SEP-2000; 2000US-0230438.
PR 08-SEP-2000; 2000US-0231242.
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PR 29-SEP-2000; 2000US-0236369.
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PR 20-OCT-2000; 2000US-0241221.
PR 20-OCT-2000; 2000US-0241785.
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PR 08-NOV-2000; 2000US-0246613.
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PR 17-NOV-2000; 2000US-0249210.

PR 17-NOV-2000; 2000US-0249211.
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PR 17-NOV-2000; 2000US-0249214.
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PR 17-NOV-2000; 2000US-0249245.
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PR 17-NOV-2000; 2000US-0249265.
PR 17-NOV-2000; 2000US-0249297.
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PR 01-DEC-2000; 2000US-0250391.
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PR 08-DEC-2000; 2000US-0251856.
PR 08-DEC-2000; 2000US-0251868.
PR 08-DEC-2000; 2000US-0251869.
PR 08-DEC-2000; 2000US-0251989.
PR 08-DEC-2000; 2000US-0251990.
PR 11-DEC-2000; 2000US-0254097.
PR 05-JAN-2001; 2001US-0259678.

XX
PA (HUMA-) HUMAN GENOME SCI INC.
XX
PI Rosen CA, Barash SC, Ruben SM;
XX
DR WPI; 2001-465570/50.
XX
XX Isolated nucleic acid molecule encoding a reproductive system antigen
PT is used in preventing, treating or ameliorating a medical condition -
XX
PS Disclosure; SEQ ID NO 6323; 1297bp + Sequence Listing: English.
XX
CC The present invention provides the protein and coding sequences of a
CC number of human reproductive system related antigens. These can be used
CC in the prevention and treatment of reproductive system disorders,
CC including cancer. The present sequence is a genomic sequence encoding a
CC protein of the invention.
XX
SQ Sequence 6063 BP; 1655 A; 1162 C; 1259 G; 1987 T; 0 other;

Query Match      24.7%; Score 427.6; DB 22; Length 6063;
Best Local Similarity 46.4%; Pred. No. 8.4e-99;
Matches 475; Conservative 211; Mismatches 298; Indels 40; Gaps 6;

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Db 3541 cattttaaaggaac--agcataaaatttgaaatttgccagccagctgtatgcagca 3598

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QY	1190	smnagcthgargtntgtgarcnccnccatltmngnttgcaraaraethaaynaaycnctga	1249
Db	3959	caagaatctgaggtcttggaaccccccactctcagattctagaagctgtcatgtacatctga	4018
QY	1250	tgccmngncaracttgcngtngngtngngnwnsnltgmgnaacnswngcmngntng	1309
Db	4019	tgctccagcgaacaaagtgtgtcccgagggccagagccttcaatgcagagaaacctgttagg	4078
QY	1310	tncaaraagtnaaygtntgngtggagctccnccncaaymngnttgcnsngtngnccnwnw	1369
Db	4079	tg-tgaaggaataatgctgggtctgtagagcccccacacagaatccctacgtgggcacaccta	4137
QY	1370	smnngcngctnmngnwnsnccnswnswnmngntncaraargtmngnwnsnacngayw	1429
Db	4138	gtgagagctgtgcgaagaagggccacgcctcttcagaccacccagatgtgatctcacgaca	4197
QY	1430	snynncaracgctnccngataarawsnacngayancarctgycarcnctngtaarngcng-cn	1488
Db	4198	gctgtatccgtgcacccctgcgaataagccacagacccttaaacgcgcgttgaaagagccaca	4257
QY	1489	gnaatggarwsngtncntctayaaracngtntgncgarcctgtaacnaaraacngtngnath	1548
Db	4258	ggagctggggactataaccctctgtagagccacagggccagagctgtgccaaagactaaggaa	4317
QY	1549	taytntyncaayctgcaycayaytngaygtcmngnagayngtncarcmngnrcaytctgyn	1608
Db	4318	tacctctgtgacatcttgcactctgcctgtagctgtagacatgtagcagggatagagatatctga	4377
QY	1609	gcnctmngntcttgayctgcnaacngntctymnagcnctatgtggncnctg-ncnynctng	1667
Db	4378	acgtataaatcttgacgcgcgcctgtgattcttgactctgacatgagggctgtgtaagcccttg	4437
QY	1668	ycttgcngcacttcttgcncnttvgnaacngcnngnttlyancarcarytntaytntcayctg	1727
Db	4438	ctgtgggcactctctcccaatttgaaatgctgatacttaccatccattacccatctatcccat	4497
QY	1728	yatg 1731	
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Qy 1688 tlygnaacngcngtntlytncacartgytlnataytncaytgyat 1730
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RESULT 10
ID AAL03634 standard; DNA: 6063 BP.
XX AAL03634:
AC 21-NOV-2001 (first entry)
DT
XX
DE Human reproductive system related antigen DNA SEQ ID NO: 6322.
XX
KM Human: reproductive system related antigen: reproductive system disorder:
cancer; gene therapy; ds.
XX
OS Homo sapiens.
XX
PN WO200155320-A2.
XX
PD 02-AUG-2001.
XX
XX 17-JAN-2001; 2001WO-US01339.
XX
PF 31-JAN-2000; 2000US-0179065.
XX 04-FEB-2000; 2000US-0180628.
PR 24-FEB-2000; 2000US-0184664.
PR 02-MAR-2000; 2000US-0186350.
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PR 01-SEP-2000; 2000US-0229344.
PR 01-SEP-2000; 2000US-0229345.
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PR 05-SEP-2000; 2000US-0229513.
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PR 14-SEP-2000; 2000US-0232397.
PR 14-SEP-2000; 2000US-0232398.
PR 14-SEP-2000; 2000US-0232399.
PR 14-SEP-2000; 2000US-0232400.
PR 14-SEP-2000; 2000US-0232401.
PR 14-SEP-2000; 2000US-0233063.
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PR 20-OCT-2000; 2000US-0241785.
PR 20-OCT-2000; 2000US-0241786.
PR 20-OCT-2000; 2000US-0241787.
PR 20-OCT-2000; 2000US-0241808.
PR 20-OCT-2000; 2000US-0241809.
PR 20-OCT-2000; 2000US-0241826.
PR 01-NOV-2000; 2000US-0244617.
PR 08-NOV-2000; 2000US-0246474.
PR 08-NOV-2000; 2000US-0246475.
PR 08-NOV-2000; 2000US-0246476.
PR 08-NOV-2000; 2000US-0246477.
PR 08-NOV-2000; 2000US-0246478.
PR 08-NOV-2000; 2000US-0246523.
PR 08-NOV-2000; 2000US-0246524.
PR 08-NOV-2000; 2000US-0246525.
PR 08-NOV-2000; 2000US-0246526.
PR 08-NOV-2000; 2000US-0246527.
PR 08-NOV-2000; 2000US-0246528.
PR 08-NOV-2000; 2000US-0246532.
PR 08-NOV-2000; 2000US-0246609.
PR 08-NOV-2000; 2000US-0246610.

XX (HYSE-) HYSEQ INC.
 PA
 XX
 PI Dimaac RT, Liu C, Tang YT;
 XX
 DR WPI: 2001-639362/73.
 DR P-PSDB: ABG21075.
 XX
 PT New isolated polynucleotide and encoded polypeptides, useful in
 PT diagnostics, forensics, gene mapping, identification of mutations
 PT responsible for genetic disorders or other traits and to assess
 PT biodiversity -
 PS
 PS Claim 1, SEQ ID No 23066; 103pp; English.
 XX
 CC The invention relates to isolated polynucleotide (I) and
 CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
 CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
 CC and gene mapping, and in recombinant production of (II). The
 CC polynucleotides are also used in diagnostics as expressed sequence tags
 CC for identifying expressed genes. (I) is useful in gene therapy techniques
 CC to restore normal activity of (II) or to treat disease states involving
 CC (II). (II) is useful for generating antibodies against it, detecting or
 CC quantitating a polypeptide in tissue, as molecular weight markers and as
 CC a food supplement. (II) and its binding partners are useful in medical
 CC imaging of sites expressing (II). (I) and (II) are useful for treating
 CC disorders involving aberrant protein expression or biological activity.
 CC The polypeptide and polynucleotide sequences have applications in
 CC diagnostics, forensics, gene mapping, identification of mutations
 CC responsible for genetic disorders or other traits to assess biodiversity
 CC and to produce other types of data and products dependent on DNA and
 CC amino acid sequences. AAS64197-AAS94564 represent novel human
 CC diagnostic coding sequences of the invention.
 CC Note: The sequence data for this patent did not appear in the printed
 CC specification, but was obtained in electronic format directly from WIPO
 CC at ftp.wipo.int/pub/published_pcl_sequences.
 CC
 SQ Sequence 1580 BP; 357 A; 412 C; 328 G; 483 T; 0 other;
 Query Match 23.1%; Score 400.4; DB 23; Length 1580;
 Best Local Similarity 42.4%; Pred. No. 2.6e-92;
 Matches 466; Conservative 215; Mismatches 383; Indels 34; Gaps 4;
 Oy 661 aaytlaytawnsnttygatyngatyncaaytgyaargynaathtgytaty 720
 Db 1107 AATTCGATACGAACAAGACTGTGGCATGTAACTAGTAACTCCGACGATCA 1048
 Oy 721 mgnaaarathylngcnaayaargarathwsnaarcarcarwnathcargatn 780
 Db 1047 GCATCCCCCACTTCTTAAGGACGACACTGATTCACGCCACCTGAGATTGACAAATAC 988
 Oy 781 acntggatynltnaargentlwsntlyatlmgngargcngarayaarwsnngar 840
 Db 987 AGATGTGTTTGAATTGGAACTTATATTAAAGGAGACAGACATAAAAAATTGCA 928
 Oy 841 aaytlaytawnsnttygatyngatyncaaytgyaargynaathtgytaty 900
 Db 927 AATCTGCGACATGACATGAAAGAAAGAAACCCATTTCGCGGAGAAATTCAG 868
 Oy 901 ytlngcngnargathylngcnaayaargarathwsnaarcarcarwnathcargatn 960
 Db 867 CTGTTGCGAAGAAATTTGATTAATACAGAGACCAATGTTAATCACCAGACACACGG 808
 Oy 961 garaayathwsnttgcacmngnwsnscarcarwnathcargatn 1020
 Db 807 AAAAATGTTTTCAGGGCCTCAGACACCTTATGCGACGCCCTCCATCCAGGCCACG 748
 Oy 1021 mgnccmngmnaa---tggatlytygngcngcngcngnwnylnhtygtynar 1077
 Db 747 AGGCTTAGAGAAATATATGCTTGTGGCTGGGCTCAGACCTTGCTCTTGTGAG 688
 Oy 1078 ccmmngaytlnctnctygtncngtlnaaysng----- 1114

Db 687 TCTCAGACTGTGTCCCTGATCCATCTGTGGCTAAAGGGCCATATACAGCTGAA 628
 Oy 1115 -cngtngcwnsngargngcwnsnccnaarcentlygcarytlnccwnsngngtngarcn 1173
 Db 627 ACCATTGCTTCAGAGGGGTAAAGCCCAAGCATGTGAGAGGCTTACATGCTGTGGACCT 568
 Oy 1174 gtlngngcnaararwsnmgathtgargtngtggarcncncnahtmgntlycaraarath 1233
 Db 567 GTGGGTACACAGAGTCAAGAACTGAGGTTTGGGAACCTGTGCTTACGATTTCAAGAGATG 508
 Oy 1234 taygnaayccnltgagtcmmngcaraartlycngtngtngtngtngtngtngtngtngtng 1293
 Db 507 TATGGAACACCTGGATATTCAGGACAGAGGTCTGCTGAGGGGTAGAGCCCTCATGTGAGA 448
 Oy 1294 acnwsngcmmngtngtncaraargnaaytngntngtngarcncncncaymngtngtncn 1353
 Db 447 ACCCTGTAGGGAATTCAGAAAGGAATGTGGGCTTGGAGCCCTCCACACAGAGTCCCC 388
 Oy 1354 wsnngngcncnwsnsmngcngtngtngtngtngtngtngtngtngtngtngtngtngtngtng 1413
 Db 387 ACTGG-----GACTAGTGAGCTGTGAGAAAGGGCCACCATCTCCAGACCCAGAT 334
 Oy 1414 gmgngwnsacngaywnytnrcarayaingtncngaraarwsnacngayacncartgycar 1473
 Db 333 GATAGACCACTGACAGCTTGATCTGCTACTGGAAGAAAGCACAAACATCTAATGCCAT 274
 Oy 1474 ccngtlnaargcngcngnargatngtngtngtncnlayaaraacngtngtngtngtngtngtng 1533
 Db 273 CCTGTGAAGACAGCCAGAGGGAGCTGTACCTGCAAAAGCCACAGGGGAGACCTTCCC 214
 Oy 1534 aaracngtngnathtlytlnrcaytygcaaytyngyngtngmngcayngngtngaar 1593
 Db 213 AAGGCTGTAAGAACCCACCTCTCATCAATGAGCTGTGAGTGAAGCTGGAGTCAAA 154
 Oy 1594 mngaycaytlytyngcngtngtngtngtngtngtngtngtngtngtngtngtngtngtng 1653
 Db 153 GGAGATCATTTTGGAGTTTAAGATTGGCTGCCCACTGATTTAGAGATTTGATGGATGGG 94
 Oy 1654 ccngt-ncenyntngtlytyngcncartlytlycncntlytyngcngcngtngtngcncart 1712
 Db 93 CCTGTAGCCCTTGTGTTTGGCAATTTCCATCTGGAATGAGTGTATATCCAAATG 34
 Oy 1713 ytlntlaytncaytgyat 1730
 Db 33 CCCGTACCCCATTTATAT 16

Search completed: July 8, 2002, 16:25:55
 Job time: 12728 sec

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Db 47624 atgtttgtcttggtggcccaaggccttctgtgtcagctcctaagacttgctcc 47683
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Oy 1095 ntgttncngtnaaywscngtngcnwngargngcnwscncaarccnggaaryt 1154
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Db 47684 cccatcc-----cagcagtgtcaaaagggccaatgtacagcttagaactt 47731
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Oy 1155 ncnwngngtngarcngtngngcnaaraarwsmngnathargtntlgarcnc 1214
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Db 47732 tgcctcagaggtgcaagc-----cc 47751
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Oy 1215 nathmgtnttcaaraarathaygnaaycntgtatgccmmncaraarttgcngtng 1274
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Db 47752 caagccttggtgtcctacatgtgtgtgtgacctgacagacacagaagttctgcact 47811
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Oy 1275 ngtnngwsmwtgtgmgcnwscngcmngtngtncaraaragnaaytngnttgg 1334
      :| | | | | | | | | | | | | | | | | |
Db 47812 ggtggaacctcctatgtgaacctctgtcagtgagcagtgataagtgaggttcgga 47871
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Oy 1335 rcncncaymngtncnswngngcnc-----cnwsmngmngcngtngmngnws 1388
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Db 47872 gcccccacacacatccccaatggcagctggcactgtactgtactgtgaactgtgagaag 47931
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Oy 1389 ncnncnswsmwtmngtncaraaragmngmwnacngaywstnrcaraaytncnca 1448
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Db 47932 gccaccatctccagaccacagaatgttagatccactgtactgtgaacctgtgcacttg 47991
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Oy 1449 raarwsmnagayacnarcarycargcngtncnargcngcngnatlgarwngtncnta 1508
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Db 47992 aaaaagccacagacactaacaacagcctgtgagagcgacgtggaagagagctgaacctg 48051
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Oy 1509 yaaraengtngtngcngarytlnacnaaraengtngnathlytntynlncaytgcaya 1568
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Db 48052 caaaacacagagacagagctgtgccaaagtcatgtgagccacactcttgacatgacctg 48111
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Oy 1569 yytngaytngtncayngtncnarmngaycaytltgngcngtltmngtnttgaatgtcc 1628
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Db 48112 ctgaaatgtgagacatgaaacaaagagatcatlttgagactttagacttgacct 48171
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Oy 1629 nacyngtngnactnlaytngcncngt-nccnyntngtltgngacarttlytncnt 1687
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Db 48172 acccggttccgactgtgatgtggcctgtgcccccttaatttgccaattatcccat 48231
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Oy 1688 tlygnacngcngtnttlyancartgytntaytncaytgyat 1730
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Db 48232 ttggaaatgggtatattaccacatgctgtaccccatctcat 48274
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RESULT 2
US-08-687-080-59/c
; Sequence 59, Application US/08687080
; Patent No. 5965427
; GENERAL INFORMATION:
; APPLICANT: Gregory Dolganov
; TITLE OF INVENTION: Human RAD50 Gene and Methods of Use Thereof
; NUMBER OF SEQUENCES: 175
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Dehlinger & Associates
; STREET: 350 Cambridge Avenue, Suite 250
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94306
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/687,080
; FILING DATE: 17-JUL-1996
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
```

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; APPLICATION NUMBER: US 08/592,126
; FILING DATE: 26-JAN-1996
; ATTORNEY/AGENT INFORMATION:
; NAME: Sholtz, Charles K.
; REGISTRATION NUMBER: 38,615
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (415) 324-0880
; TELEFAX: (415) 324-0960
; INFORMATION FOR SEQ ID NO: 59:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 14855 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; HYPOTHEICAL: NO
; ANTI-SENSE: NO
; ORIGINAL SOURCE:
; INDIVIDUAL ISOLATE: 5' END OF INTRON 2 OF RAD50 GENOMIC
; INDIVIDUAL ISOLATE: SEQUENCE
; US-08-687-080-59
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Query Match 10.3%; Score 177.6; DB 2; Length 14855;
Best Local Similarity 44.4%; Pred. No. 2.2e-39;
Matches 221; Conservative 90; Mismatches 172; Indels 15; Gaps 4;
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Oy 1244 cntgatccmncnrcaraarttgcngtngngtngswmwtltgmgcnwscngcm 1303
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Db 14854 CCTGATGCCCCAGGCAAGATGTTGCTCAGGGGGGGCCCTTATGGAANAACCTCTCTA 14795
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Oy 1304 gngtngtncaraaragaytngntltggarcncncaymngtntncnswngngcnc 1363
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Db 14794 GGGCAATATGGAAGGAATGTGGGTTGAACCCAC--AGAATTCTATGAGGGGAC 14737
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Oy 1364 cnwsmnngcngtngtngmngwnccncnswsmngntl-----ncaraar 1413
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Db 14736 TGCTAGTGAGCTGTGAGAAAGACGACCTGTCTCAGACTGTGATGCCAGAAAT 14677
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Oy 1414 gnmngnswngaywstnrcaraaytncnrcaraarwsmnagayacnarttgycar 1473
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Db 14676 AATAGATCCACTGACAGCTTGACCTGTGACCTGGAAAACTGACGACACTCAACACAG 14617
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Oy 1474 cngtncnargcngngnatlgarwngtncnlayaaraengtngcngarytlnacn 1533
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Db 14616 CCTGTGAANAACAGCCAGAGAGAGGTATACCTGCAAAAGCC--AGAAATGAGACTGCC 14559
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Oy 1534 aaraengtngnathaytntaytncaytgcayaytngaytngmngcngngtncnaar 1593
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Db 14558 AAGCCATGGAAGCCACCTCTTCATCAGATGACCTGTGATGTGAGACATGAGTCAAA 14499
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Oy 1594 mngaycaytngngcngtngmngtltgaytgcacnngnttymnactnlayatggn 1653
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Db 14498 GGAGATATTTCTGAGAGCTTTAAGATACACTGCCCACTGAATTTGGACTTGACAGCGGG 14439
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Oy 1654 cngt-nccnyntngtltgngcncarttlytncnttlygngacngcngtnttlyancartg 1712
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Db 14438 CCTGTAGCCCTGTTGTTGGCCAAATTCCTCCATTTGGAATGCTGTATTTGCCCAATG 14379
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Oy 1713 yytntaytncaytgyat 1730
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Db 14378 CCTGTATCCCATTTGAT 14361
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RESULT 3
US-08-991-789A-29
; Sequence 29, Application US/08991789A
; Patent No. 6225054
; GENERAL INFORMATION:
; APPLICANT: Fridakis, Tony N.
; Smith, John M.
; Reed, Steven G.
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OY 1070 gysltncarcmmgnagaylntgntcncttgytncctcgtlnaaaywsngcngtngcnswngarg 1129
Db 676 gayatlgcngcnynltbgcaywsnaartaaygarwsngayaenylngarcargayltingar 735
OY 1130 gngcnwsmcnbaarcscntgycarytncnswsnhgntngarcngtngngcnaararw 1189
Db 736 mgnyntltcargayrltmngcnctylnatayltnaaaycncayactaygtlmgmngcgn 795
OY 1190 snmgatnhtgargtntggarcncnccnathmgntlycaraaathlaygnaaycnclyga 1249
Db 796 ylncaymncaytlaygncngcngarylnathgayltnmgnggncnccnathcngcncayltn 855
OY 1250 lgcncmgncaararctlygcngtngngcngtngngnswntslgmgnaacnswngcngmngtng 1309
Db 856 ylmgngngaraayacnlytngcncarcsntlgygtlnaayathyltngaycngtntynctnly 915
OY 1310 tncaraargnaaygltngnttggarcncnccncaaymgngtncnswngngcncnswanw 1369
Db 916 yltnaararathcncngargayltnacnaarathalgaargltnarcayltggarcngar 975
OY 1370 snmgcngcgtlmgmngnswncncnswsnsmgnyltncaraargmngmgnwnaengayw 1429
Db 976 aarylnatgytngargargarcnactlytlyacntayltnghytlngcnyltnccnccn 1035
OY 1430 snyltncarcayltnccngaraararwsnagayacnartgycarcngtlnaargcngcng 1489
Db 1036 gncncncnswntlytlggaararayltnatgyltnatgmgnccncaengaygmgngargtln 1095
OY 1490 gnatg 1494
Db 1096 garlg 1100

RESULT 13
US-09-351-414-3
; Sequence 3, Application US/09351414
; Patent No. 6265199
; GENERAL INFORMATION:
; APPLICANT: Sheppard, Paul O.
; APPLICANT: Balndur, Nand
; APPLICANT: Delsher, Theresa A.
; APPLICANT: Bishop, Paul D.
; TITLE OF INVENTION: DISINTEGRIN HOMOLOG
; FILE REFERENCE: 98-29
; CURRENT APPLICATION NUMBER: US/09/351.414
; CURRENT FILING DATE: 1999-07-09
; NUMBER OF SEQ ID NOS: 13
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 3
; LENGTH: 2088
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: zdlntl amino acid degenerate sequence
; FEATURE:
; NAME/KEY: variation
; LOCATION: (1)..(2088)
; OTHER INFORMATION: n is any nucleotide
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)..(2088)
; OTHER INFORMATION: n = A,T,C or G
US-09-351-414-3

Query Match 2.8%; Score 48.8; DB 4; Length 2088;
Best Local Similarity 11.8%; Pred. No. 0.0026;
Matches 140; Conservative 286; Mismatches 749; Indels 8; Gaps 3;

OY 544 aaryltnwsgnaaaryltnccnynlncnctlyaaarcnathathtlyacngngtlnylnay 603
I::: I::: I::: I::: I::: I::: I::: I::: I::: I::: I::: I::: I::: I::: I::: I:::
Db 731 aratyltncaygartlywsnaarlaymgnamgcnathaaarcacaycngaygscngtnc 790

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QY	604	aaygcncamngnagaytlnaaragcnaatcvgngntltcvgntlcygmngntnccngnaay	663
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QY	664	laytlaywsnwtlttgyatngaytlnacayctgyaaygtlnaaytlbtgyltnagln	723
Db	851	tnltgysnmngnacmmgngngntngngtlnaaygarctaygntlnccnatcvgcngtncnc	910
QY	724	aaarcaalhtyngcnaaayargarathtwsnaarcarcarwsnatlhcargatlnaon	783
Db	911	argtlnylnswncarwsnlytngcncaraaytlnagnatlhcarggarcconswnsnmnga	970
QY	784	tgggtntynytlnaarngnttysnltlyatltmgngatgcngarctayaatwsnswngaraay	843
Db	971	arccnaarttgyayltgyacngarwsntltgggngngtltgyltha tggargarcngngntnw	1030
QY	844	yltnacyscnngayaygtlnaahaaaraaaraayconltysngargngaatltyaaytn	903
Db	1031	snccaywsnmngnaartltcwsnaarttgywsnathtyngarctayngaytltlytlnccmng	1090
QY	904	gncngnagarathtgyathltgyaayargaratylnaaygtlnaayccnccargayayagngar	963
Db	1091	gngngngngcngtlytntltayaymgncnacaaraytntltgarccnccnccngarttgygna	1150
QY	964	aayathswntltgcncnttgcarmgnswnsnccarcarwsnathtarwsnyltngcngtmgn	1023
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Db	1209	nyltntgytgyaaraarttgywsnlytnswnaaygngcncaytgywsng--aygngcnyt	1267
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QY	1204	tgggarcnccnatltmgntltcaraanathltaygnaayccnttggatccmmngncaraar	1263
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Db	1508	gyltaygataaytlnaayaacngargngarcngaraargna----aytgygnaaargayg	1562
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QY	1564	caygaytngaygtngmgncayggngtlnaarimgngaycaytlygngcngtymngnttgyay	1623
Db	1743	ytngtngnclaytngarjaygngnacnccnttgyggncnswnsnagatctgytngatmgnaa	1802
QY	1624	tgyccnccnccngnttlymgnaecnatlayatggcngcngtncnnytnlttgytngcncarttlyt	1683
Db	1803	rttgyttnccaratlhcargcnytnaayatlbtgwsnwtltgycnlytngaywsnaarngnaagt	1862
QY	1684	ccnttlyggnacngcngntlttcaocncarttgytntataytlncaayt	1726

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Db 554 AGCATTCAGAGGTAAGTGGTGTGTTAAAGCATTGTATTAAGGAGGAGG 613
Qy 826 cayaarsnwsnrgaaynncaycngayaytnathaaaraaraaycnclywsn 885
   ||:||||:||||:||||:||||:||||:||||:||||:||||:||||:||||:
Db 614 CATAAAGTTTGGAAATTTTGCAGCTGACATGCAATAGAAAAGAAATCCATTATCT 673
Qy 886 gargnaartlyaarlyngcngcngarathlyathgyaaygararylnaay 945
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Qy 946 cncararayaayngngaraayathwsntgacntgyacarnwsnscncarcatwsnath 1005
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Db 734 CCCCAGACAAATGGGAAATGCTCCAGGGCAGGAGAGGCTTCCACAGACCCCTCC 793
Qy 1006 aarsytnngcngtgmngnccmngmng---naartgytlygygnacngcngmngsn 1062
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Db 794 CCATCAGAGGCTTCAGGCTCAGAGAGGAAAGATGATCTCTTGGGCCAGGCCAGGGTCC 853
Qy 1063 ytnlytlygtlncarcnmngaylytngtncntlygtlncngtnaaywsngc----- 1115
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Qy 1116 -----ngtngcnsngargngcngcnsncnaarccntg 1149
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Db 914 GAAAGGGCCCAACATAGAGCTCGATTTGCTTCAGAGGGTGCAAGCTGTAAACCTTGG 973
Qy 1150 carytncnswngngtngargcngtngngcnaaraarwsnmnatlyagntlyggar 1209
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Qy 1210 cncncaathmgntlycaraarathlaygnaaycngtgyatgcnmngncaraarctlygc 1269
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Qy 1270 gtnngngtngnwsnwsntlygmngnacnswngcmngngtngtncaraaragntlygng 1329
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Qy 1330 tgggarccnccncaymngtgcnwsngngcncnswsnsmngcngtngmngmngsn 1389
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Qy 1630 acngngtlymngnactaytlytlygngcngt-nccnytnlytlytlytlytlytlytlytlyt 1688
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Qy 1689 ygnacngcngtntlytlytlytlytlytlytlytlytlytlytlytlytlytlytlytlytlyt 1730
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Db 1510 TGGAACTGCTGTATTACACAAATACCTGTACCCCATTTGTAT 1551
LOCUS BC004496 2009 bp mRNA linear HTC 12-JUL-2001
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DEFINITION Homo sapiens. Similar to hypothetical protein FLJ14058, clone
IMAGE:3831313, mRNA.
ACCESSION BC004496
VERSION BC004496.1 GI:14709139
KEYWORDS HTC.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 2009)
AUTHORS Strausberg, R.
TITLE Direct Submission
JOURNAL Submitted (12-MAR-2001) National Institutes of Health, Mammalian
Gene Collection (MGC), Cancer Genomics Office, National Cancer
Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590,
USA
REMARK
COMMENT NIH-MGC Project URL: http://mgc.nci.nih.gov
Contact: MGC help desk
Email: gcaps-remail.nih.gov
Tissue Procurement: ATCC/DCMP/DMP
cDNA Library Preparation: Rubin Laboratory
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)
DNA Sequencing by: Institute for Systems Biology
http://www.systemsbiology.org
contact: amadansystemsbiology.org
Anup Madan, Rachel Dickhoff, Jessica Fahey, Stephanie Ford, Julia
Greene, Mark Kertelman and Anuradha Madan
Clone distribution: MGC clone distribution information can be found
through the I.M.A.G.E. Consortium/LNL at: http://image.lnl.gov
Series: IRAL Plate: 14 Row: d Column: 7
This clone was selected for full length sequencing because it
passed the following selection criteria: Hexamer frequency ORF
analysis
This clone has the following problem: frame shifted.
location/Qualifiers
1. 2009
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/db_xref="taxon:9606"
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/lab_host="DH10B-R"
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BASE COUNT 556 a 429 c 513 g 511 t
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Best Local Similarity 45.2%; Pred. No. 1.1e-92;
Matches 452; Conservative 214; Mismatches 321; Indels 13; Gaps 3;
Qy 744 rgargarathwsnaarcarcarwsnathcargarytnacnsgtngnytnaargcnt 803
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Db 549 CAGATTTTAAGGGAAGCAAGACATTAAGAAATTTGCAAGAAATTTGCAAGCTGCAATGTGAT 608
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Db 669 TAACAAGAAAGCGCATGTGTAATCCCAAGACAGCAAGCAAGACATCTCCAGAGCATATCA 728
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Qy 1044 ygnacngcngcngnswnynt-----gylgyltncaarcnmngngaytngtnc--- 1094
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Db	789	TGACCTGTGCAAGCCTAGGTACTTGGCAACCTGTGTCCAGCTACTCCAGTATGTGGCTCA	848
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Db	849	AAGGGGCCAAATACAGCTTACGCATGGCTTCAGAGGGGTGCAAAGCCAAAGCTTGGCA	908
Oy	1152	rytlnccmsngnglncgarcnglncgnglnaaraarawsmgnaahgaryltblggarc	1211
Db	909	GCTTCCGATGCTGTGTGAAGCTTCGCAAAATGACAGAGCTCAAAATTTGGATTTGGGAACC	968
Oy	1212	nccnatlmgnltcyaraaathlaygnaaycngtggatgcmmncaraatlygngt	1271
Db	969	TCTGCTAGATTTTCAGAGATGTACGGAAAGTCTCCGATGCGCCAGCAGCAAAATTTGCTGC	1028
Oy	1272	ngngltnngnswanltygmagnacwmsngcmmnglntlncaaraargnaayltnnglntg	1331
Db	1029	AGGGCAGAGGCTTCATGGAGAACTCTGCTGAGGACAGTGTGAAGGAAACATGGGGTTC	1088
Oy	1332	ggarcncncncaymngltnccmsnrgngngncncmsnsmngncgtlmgmgmsncc	1391
Db	1089	TGAGCCCCACACAAATCCCTACGAGGACACACCTATGTGAGCTGTGTAGAAAAGGCC	1148
Oy	1392	ncnmsnsmngnylncaaraargmgmsnmgnaaywnynlncaraytblncngaraa	1451
Db	1149	ACCATCTCCACACCCCAATGATAGATCCACCAACAGCTTGCAACCATGCACTGGAAA	1208
Oy	1452	twsmnagnayacncarltygarccnglnaarycngngnaltlgarwslnglncnltayaa	1511
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Oy	1512	raeinglntngcngarylnachnaaraecnglmgnaahlayltnlncayltycaygayt	1571
Db	1269	AGCCACAGGGGAGGCTTCCCAACATATGGGAACCTACCTTTACACACAGAGACT	1328
Oy	1572	ngayltnmgncaygnglnaarmngaycayctlygngcnyltnmgltlygayltycnaac	1631
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Oy	1632	ngnaltymgnaactlayatlggnc--nglncnlyltnlyltylgngcaarltlyltycscntlyg	1690
Db	1389	TGGATTTACAGACTGTCATAGGGCCCTACAACCCCTTTGTTTGGCGATTTCTCCAAATTG	1448
Oy	1691	gnacngcnglntlycncarltgylnltnltnlncayltyat	1730
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/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone_lib="human B lymphocyte cdna library"
/tissue_type="bone marrow"
BASE COUNT      380 a      341 c      398 g      395 t
ORIGIN

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Query Match	23.0%	Score 398	DB 10	Length 1514
Best Local Similarity	47.1%	Pred No. 1	1e-80	
Matches 421	Conservative 182	Mismatches 256	Indels 41	Gaps 4

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RESULT      3
LOCUS       BE512633
DEFINITION  BE512633      1514 bp      mRNA      linear      EST 11-OCT-2001
              32-1514 human B lymphocyte CDNA library Homo sapiens CDNA, mRNA
              sequence.
ACCESSION   BE512633
VERSION     BE512633.1
KEYWORDS    GI:16041645
SOURCE      EST.
            human.
ORGANISM    Homo sapiens
             Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
             Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE   1 (bases 1 to 1514)
            Lu,X., Cui,L. and Li,Y.
            DDRT-PCR from B cell
            Unpublished (2000)
COMMENT     Contact: xingwu lu, liangxian cui, yonghai li
            Department of Biochemistry
            Institute of Basic Medical Science, Peking University Medical College
            Dongdan Sanjiao 5, Beijing, P.R.C, 100005
            Tel.: 86-010-65296951
            Email: luxingwu@263.net
            full cDNA sequence.
FEATURES             location/Qualifiers
             1..1514

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QY	862	abhaaraaraaayacntlytynsngairgnaaarlttyaarlytngcngnagathty	921
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QY	922	tyaaaygaratyltaaygtlnaaycncaragyaayggngaraayaatlhstnlygacntgy	981
Db	61	TCTAACAGAGATTAAATGTTTAAATCCCAAGACAAATGGGAAAATGTCTCCAGGCAATGC	120
QY	982	carmgwnswnsrcaarcatwsnahtaaarsnylncnlygmngcncmngm--naatfgy	1038
Db	121	CAGAGGTCCTTCACACAGAGCCCTCCCATCACAGGCGCTCAGGCTTAGAGAGAAAAATGA	180
QY	1039	tytyygnacngncngnmgwnsnylntygytynearcncmngaytlynglntcngty	1098
Db	181	TCCTCTGGGCCAGGGCCAGGGTCCCATGCTGTGTCAGGCTGGGAGCTGTGGTCTGTGT	240
QY	1099	gtncnqntlaaaywsnrg-----nglncnsmn	1125
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QY	1186	aarwsnmnathtgarlytntbgyarccnccnathmgtlycraaraatlhaygnaaycn	1245
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QY	1246	tygatlycncmgnrcaratltycngltnngltnngnswnslygmngacnwsngcnn	1305
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QY	1366	wsnwsnmngncngltnmgmgnwsncncncnswnsnmngylncraaraaygmngnswncn	1425
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QY	1426	gawywnlyncarayltnccngaraarwsnaengayaencarltgyarcngltnaarycn	1485
Db	601	GACACCTTGCTCCATGACACTGTGAAAAACCGCAGA-----CAACACACACCCATGAAAACA	656
QY	1486	gengnatlygarwsngltnccnlayaaracongltnngltnearcngltnaarnaaracngltn	1545
Db	657	GCTAGGACGAGAGGCTGTACCTGCAAAAGCCACAGGGGGGAGGCTGCCAAGATCATTTGGGA	716
QY	1546	athlaytynlncayltygcaygayylnaaygltnmgncaygngltnaarmngnayaclty	1605
Db	717	ACCCACCTCTTCATCACCATGACTGTGATGGAATTTGGATTAAAGGAGATCATATTTT	776
QY	1606	gngcnylnmgnltygayltygcnaacngltnmgnaactlayatbgygncngt--ncnylt	1664
Db	777	GGAGCTTTAAGATTTGATGCTGCCCTCTGTGATTTTCAGACTTGCATGCGGGCGTGAAGCCCT	836
QY	1665	ngaytlygncartlytlytlycnclytynagncngltnlycencarlttyltnlaytlnca	1724
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Db      897  TTGTAT 902

RESULT  4
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LOCUS   603021014F1 NIH_MGC_114 Homo sapiens cDNA clone IMAGE:5191803 5',
DEFINITION
mRNA sequence.
ACCESSION BI48505.1 GI:15327733
VERSION   BI48505
KEYWORDS  EST.
SOURCE    human.
ORGANISM  Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 870)
AUTHORS  NIH-MGC http://mgc.ncl.nih.gov/.
TITLE     National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL   Unpublished (1999)
COMMENT   Contact: Robert Strausberg, Ph.D.
          Email: cgaabs-remail.nih.gov
          Tissue Procurement: Life Technologies, Inc.
          cDNA Library Preparation: Life Technologies, Inc.
          cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)
          DNA Sequencing by: Incyte Genomics, Inc.
          Clone distribution: MGC clone distribution information can be
          found through the I.M.A.G.E. Consortium/LNL at:
          http://image.lnl.gov
          Plate: LLM11479 row: k column: 04
          High quality sequence stop: 849.
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/db_xref="taxon:9606"
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/clone_1id="NIH_MGC_114"
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/note="Organ: brain; Vector: pCMV-Sport6; Site:1: NCL;
Site:2: EcoRV (destroyed); RNA source anonymous pool of 6
male brains, age range 23-27 yo. Library is oligo-dT
primed and directionally cloned (EcoRV site is destroyed
upon cloning). Average insert size 1.5 kb, insert size
range 1-3 kb. Library is normalized and enriched for
full-length clones and was constructed by C. Gruber
(invitrogen). Research Genetics tracking code 019. Note:
this is a NIH_MGC Library."
BASE COUNT 222 a 211 c 256 g 181 t
ORIGIN

Query Match 19.5%, Score 336.8; DB 10; Length 870;
Best Local Similarity 44.6%, Pred. NO. 1e-66;
Matches 378; Conservative 161; Mismatches 273; Indels 35; Gaps 3;

Db      847  caycnggaaayaygnatnaaraaraaycnyntlyngargnaattlyaytngcn 906
2  CAGCTGTATGATGAAGTAAGTAAGTAAGTAAGTAAGTAAGTAAGTAAGTAAGTAAGT 61
907  gongaratlygathlygaaygargarylnaayglnaayccnccargayaaaygngaraya 966
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967  atlwentggaactcycaarmgwnsnwncarwaarwnatnaarwnytnngntgmgnc-c 1025
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Db      122  GTCTCCGGGCCATGTGCAGAGACCTTCATGGCGCCCTCCCATTCACAGGCTGGAGCCAG 181
1026  nmngnnaarbtgtytygagncngcngnwnytnlytygtnncarccmgnga 1085
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1086  ytngtncntlygtncngt-----naayws 1112
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Db      242  CTTGTGCCCTGTGTCCAGTCGCTCCAGCTGTGACTGAAAGGGGCCAAGTAGAGCTCG 301
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Db      302  GGGTGTGGCTTTCAGAGGGGTGGAGCTTAACTTAACTTAACTTAACTTAACTTAACTTAA 361
1173  ngtnngcnaaraarwaarwnatnaarwtgargtnltggagccnccnaatnngntlycraarat 1232
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Db      362  TACAGGTGCACAGAAAGTCAGAACTGAGGTTTGGGAACTCTGCTTGATTTTCAAGAGAT 421
1233  hlaygnaaycncntlygaatgcnmgncnaararltlycngtngngtngtngnswnsn 1292
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Db      422  GTTTGGAATGCTGTGATGCGCCAGCAAAAGTTGTGTCAGAGAGTGGGCCCTTATGAGAG 481
1293  nacnswngcmngtngtngtncaraaragayngtngnttggarccnccnaayngtncnc 1352
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Db      482  AACCTCTGTGGACAGTGCAGAGGAAAGTGGGATCGGAGCCCGCAGAGAGATGCCA 541
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1413  rgnmgwnsnacngaywsnylnccaragylncngararwaarwnacngayacnartlyca 1472
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722  CAGACCATGTGGGAACCCACTTGTGATCAGCGGTGACCTTGATGCGAGACCTGGAGTCA 781
1593  rmgngaycaatlytngcngnt-nmngtlygalytgcnaacngntlymgnaactayatlg 1651
782  AGGAGATCATTTTGGAGCTTAAACATGTGACCTGCTGATTCAGACTTGATTCAGATTG 841
1652  gncncgt 1658
842  GGGCCT 848

RESULT  5
LOCUS   BM472108 1050 bp  mRNA  linear  EST 05-FEB-2002
DEFINITION
AGENCOURT_6465359 NIH_MGC_72 Homo sapiens cDNA clone IMAGE:5539381
5', mRNA sequence.
ACCESSION BM472108
VERSION   BM472108.1 GI:18521150
KEYWORDS  EST.
SOURCE    human.
ORGANISM  Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 1050)
AUTHORS  NIH-MGC http://mgc.ncl.nih.gov/.
TITLE     National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL   Unpublished (1999)
COMMENT   Contact: Robert Strausberg, Ph.D.
          Email: cgaabs-remail.nih.gov
          Tissue Procurement: ATCC/DCPD/DRP
          cDNA Library Preparation: Life Technologies, Inc.
          cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)
          DNA Sequencing by: Agencourt Bioscience Corporation
          Clone distribution: MGC clone distribution information can be
          found through the I.M.A.G.E. Consortium/LNL at:
          http://image.lnl.gov
          Plate: LLM1223 row: m column: 14
          High quality sequence stop: 658.
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Source
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/organism="Homo sapiens"
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Qy 1158 nwgngngtngatrcngtngngngncaaraarwsmgmacthargtntggarcnccnat 1217
Db 142 ATGTGTCTTGGTCTGTGAGTCACAGAGA-----ATCGGGCTTTGGAACTCCACT 136
Qy 1218 tmgntctaraarathayagaaayocnttgatgctcmgmcaaraarttgcngtngngnt 1277
Db 197 TAGATTTCAGAAAGATGATGATAAATACCTGATGCTCCATGCAAAAGTTCTGTAGGGTC 256
Qy 1278 ngmnmwsmnltgmgngnagmngnngnngtngtncaraaragngaytngnttggarc 1337
Db 257 AGGGCCCTCATGAGAACTCTCTCTAGGGCACTGAGGAAAGGAAATGTGAGGTCAAGGC 316
Qy 1338 nccngaymgngtngcnwngngngcnwsmngmngcngtngmngmngmngmng 1397
Db 317 CCACACAGAGTCCCTACTAGGGCCACCAGCTGAGACTGTGAGAGAGGCGCACTGTC 376
Qy 1398 nwmngnylncaraaragmngmngmngmngmngmngmngmngmngmngmngmng 1457
Db 377 CTCAGACTCCAGAAATGTGATGATCCATGCTTCCATCATGCTCTGCAAAAGGCAC 436
Qy 1458 ngayacnartgycartcngtngnagngngnagatggagawngtngnctayaraagnt 1517
Db 437 AGACACTCAATGCCAGCCCATGAAAGCAGCGGAGAGGCGTGTACCTCAAGGCCAC 496
Qy 1518 ngtngngartl-nacnaaracngtngncthayaayltncaaytgycaaytngay 1576
Db 497 AGGGCGAGGCTGTCCAAAGCATGAGACCCCTTCCCTGCTGCTGAGTGTG 556
Qy 1577 tmgngcaygngtngnagngaycaytlygngngmngtngmngtngaytngay 1636
Db 557 TGAGATATGAGTCAAGAGATCATGTGAGCTTGAATTTGACTGTGCTGAT 616
Qy 1637 tmgngactlayatgngcngt-nccnylntgyltngcngarttlycngtlygngn 1695
Db 617 TTGGACTTGCATGGGGCTGAGCTCTTGTGTTGGCAATTTCTCCATTTTGATG 676
Qy 1696 gcngtntlyacna 1709
Db 677 GCTGTATTACCA 690

RESULT 13
AG116938
LOCUS AG116938 720 bp DNA linear GSS 03-NOV-2001
DEFINITION Pan troglodytes DNA, clone: PTB-124K01.R, genomic survey sequence.
ACCESSION AG116938.1 GI:16737457
VERSION GSS: GSS (genome survey sequence).
KEYWORDS Pan troglodytes male lymphoblast DNA.
SOURCE Pan troglodytes
ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Pan.
TITLE Fujiyama, A., Hattori, M., Toyoda, A., Taylor, T. D., Yada, T.,
JOURNAL BAC end sequences of library PTB
REFERENCE Unpublished
AUTHORS Fujiyama, A., Hattori, M., Toyoda, A., Taylor, T. D., Yada, T.,
TITLE Fujiyama, A., Hattori, M., Toyoda, A., Taylor, T. D., Yada, T.,
JOURNAL Direct Submission
REFERENCE Submitted (02-AUG-2001) Aaso Fujiyama, The Institute of Physical
and Chemical Research (RIKEN), Genomic Sciences Center (GSC);
1-7-22, Suenho-chou, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
Tel: 81-45-503-9111, Fax: 81-45-503-9170
Clones are derived from the chimpanzee BAC library PTB this BAC end
was generated during the Rtd process and may have higher chance of
clone tracking errors.
PRIMERS
Sequencing: M13Rev

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LIBRARY
Vector : pKS145
R.Site 1 : SacI
R.Site 2 : SacI.
Location/Qualifiers
1..720
/organism="Pan troglodytes"
/db_xref="taxon:9598"
/clone="PTB-124K01.R"
/sex="male"
/cell_type="lymphoblast"
/clone_1lb="PTB Chimpanzee Male BAC library"
BASE COUNT 181 a 186 c 187 g 165 t 1 others
ORIGIN

Query Match
Best Local Similarity 16.5%; Score 286.2; DB 12; Length 720;
Matches 298; Conservative 112; Mismatches 200; Indels 4; Gaps 3;

Qy 1114 gongtngcmwngatrgngngcmwngnccnarttgaytngcnwngngtngarcn 1173
Db 110 GCCATGCTTCAGATGTCAGCAAGCCAGATGCTTGTGCTTCCATGTCATTTGAGCT 169
Qy 1174 gtlngngcnaaraarwsmngmngnathargtnttgarcnccnaltmgntctaraarath 1233
Db 170 GCGGTCACAGAAATGCAAGATTTGAGGTTTGAAACCTCCACCTCAATTTCAAGACATG 229
Qy 1234 taygnaaycngtnggagcngmngncaraarttlycngtngngtngngmngmngntgmn 1293
Db 230 TATGCAAAATGCTGATGATGCCAGGAGAGATTTCTGCAAGGGAGTGGCTCTCATGAGAA 289
Qy 1294 acnwsngmngtngtngcaraaragngaytngngtggagacnccnccaymgngtngn 1353
Db 290 ACCTTCTAGGCGAGTGCAGAGAGAAATGTGGCTCGAGCCGCCACACAGGTCCT 349
Qy 1354 wngngngcnccnsmwsmngmngngtngmngmngnccnccnsmngnylncaraar 1413
Db 350 ACTGGCG-ATCACCTTAAGAGAGCTGTGAGAGACAGCCAGCTCTTCCAGACCCAGAT 408
Qy 1414 gmgmngmngnagaywsmnylncarcaytngcngnagaaarwsmngayacnartgyoar 1473
Db 409 GCTAGATCCACTGCAACTTGC--ACCTGTGCTGAAAGCTGCAAGTCAATCAACATCAG 466
Qy 1474 cngtngnagngngnagatggagwngtngcngtngayaraengtngtngngarttngn 1533
Db 467 CCATGAAAGGAGCTGGAGGAGGCTGTACCTGCAAAAGACAGAGAGTGGACTTCC 526
Qy 1534 aarengtngnathaytlytngaytgycaaytngaytngmngcayngngtngaar 1593
Db 527 GAGACATGGGAACCTTCTCTCAACAGGTCAGACTGAGTGTGACATGGAATTCAAA 586
Qy 1594 mngaycaytlygngngtngnttlygaglygcnaengnttngncaatlayatggg 1653
Db 587 AGAGATCATATGAGGCTTTAATTTGACACCCACTGATTTGGACTTGCATGCG 646
Qy 1654 cngt-nccnylntgyltngcngarttlycngtlygngcngngtngtngayacartg 1712
Db 647 CCTGTAGCCCTTTGTTTGGCAATTTCTCCATTGGAATGTTGATTACCTAATG 706
Qy 1713 ytlngaytngcayt 1726
Db 707 TCTGTATTCCACTT 720

RESULT 14
AQ037711
LOCUS AQ037711 623 bp DNA linear GSS 11-JUL-1998
DEFINITION CIT-HSP-2337G10.TR CIT-HSP Homo sapiens genomic clone 2337G10. DNA
ACCESSION AQ037711
VERSION AQ037711.1 GI:3303543
KEYWORDS GSS.

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SOURCE      human.
ORGANISM    Homo sapiens
REFERENCE   Adams,M.D., Rounsley,S.D., Zhao,S., Field,C.E., Bass,S., Linher,K.,
AUTHORS     Golden,K., Berry,K., Granger,D., Suh,E., Wible,C., Shizuya,H.,
            Simon,M. and Venter,J.C.
TITLE       Use of a random BAC End Sequence Database for Sequence-Ready Map
            Building (1998)
JOURNAL     Unpublished (1998)
COMMENT     Other-GSSs: CIT-HSP-2337610.TF
            Contact: Mark Adams
            Department of Eukaryotic Genomics
            The Institute for Genomic Research
            9712 Medical Center Dr., Rockville, MD 20850, USA
            Tel: 301 838 0200
            Fax: 301 838 0208
            Email: mdamas@tigr.org
            Clones are available from Research Genetics (info@resgen.com). BAC
            end search page:
            http://www.tigr.org/tdb/humgen/bac_end_search/bac_end_search.html.
            Seq primer: M13 Reverse
            Class: BAC ends.
FEATURES    location/Qualifiers
            source      1. 623
                        /organism="Homo sapiens"
                        /db_xref="taxon:9606"
                        /clone="2337610"
                        /clone_lib="CIT-HSP"
                        /sex="Male"
                        /cell_type="Sperm"
                        /note="Vector: pBelobAC11; site_1: HindIII; site_2:
                        HindIII"
BASE COUNT  160 a      146 c      172 g      145 t
ORIGIN
Query Match      16.4%; Score 283.8; DB 12; Length 623;
Best Local Similarity 48.1%; Pred. No. 1.5e-54;
Matches 286; Conservative 111; Mismatches 196; Indels 2; Gaps 2;

QY 1138 cnaarccnltgagcarylncnswngnltngarcngltngngcnaarararwsmgnath 1197
      |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 26 CTCAGAGCTTTGGCAGCTTCCACATGCTTGAGCTCGGGGTACACCAAGTCAAGAACT 85

QY 1198 gaggltngt-ggarcncnathngntlycaraanathtaygnaaycngtgcmmg 1236
      |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 86 GAGCTTGTGAGAGACCTCCACCTACATTTTCAGAAAGATGATGGAATGCTTGATGTCAG 145

QY 1257 ncaaatctgcnngtngngtngnswngntgmgncmngcmmngntngncaraa 1316
      |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 146 GCAGAAATTTTGTCTCAGAGGGGAGGGCCAGATGAGAACTCTCTAGGGCAGTGGGAA 205

QY 1317 rgnaaaygtngnltgagarcncncncaymngnltncnswngnngcncnswsmngnc 1376
      |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 206 GGGAAATGTGGGCTTGAGAGCGCTCAGACAGAGCTCTACAGGGCAGCTAGTGGAGC 265

QY 1377 ngtnmgmngwnscncncnswsmngnylncaraargmngmwnscnagaywnylnc 1436
      |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 266 TGTGAGAGAGGGGCGCATTTCTCCAGACCCCTGAAATGTGATGATCCCGACACTTGCA 325

QY 1437 rcaygtlncngaraarararararararararararararararararararar 1496
      |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 326 TTATGCAACTGGAGAAAGCCACAGACACTCAACACAGCCCATGAGAAACAGCAGAGGGG 385

QY 1497 rwsngtncnltayaraarngltngnngaryltnacnaararararararararar 1556
      |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 386 AGCCATACCTGCAAGAACCCACAGAGGAGAGTGGCTGAGAGCCATGGGAACCCACCTTT 445

QY 1557 ncaytgycaaytngaytngmncaygngnltlaarmngnagaytlytgngcnyltmg 1616
      |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 446 GCATACAGCTGACCCAGATGTGACATGTGTCAAGAGATCATTTTGTGAGCTTTAAG 505

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QY 1617 ntygaytgcnaacngntlymgnacnlayatgngcngt-ncnylntgltgngc 1675
      |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 506 ATTGAGACGCCCTGCTGATTTTCAGACTGTGAAGGGGACATACCCCTTTCTTTGGCC 565

QY 1676 arttytlycnclytgnacngnltlyacncarctylyntaytlycaytlyat 1730
      |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 566 AATTTCCTCCATTTTGAATGAGCTGTATTTACCAATGCTGTATTTCCATGTGAT 620

RESULT 15
AG097258      677 bp      DNA      linear      GSS 03-NOV-2001
LOCUS      Pan troglodytes DNA, clone: PTB-098001.R, genomic survey sequence.
DEFINITION      AG097258
ACCESSION      AG097258
VERSION      AG097258.1 GI:16717775
KEYWORDS      GSS; GSS (genome survey sequence).
SOURCE      Pan troglodytes male lymphoblast DNA, clone_lib:PTB Chimpanzee Male
            BAC library clone:PTB-098001.R.
ORGANISM      Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Pan.
REFERENCE      1 (sites)
AUTHORS      Fujiyama,A., Hattori,M., Toyoda,A., Taylor,T.D., Yada,T.,
            Totoki,Y., Watanabe,H. and Sakaki,Y.
TITLE       BAC end sequences of library PTB
JOURNAL     Unpublished
REFERENCE   2 (bases 1 to 677)
AUTHORS      Fujiyama,A., Hattori,M., Toyoda,A., Taylor,T.D., Yada,T.,
            Totoki,Y., Watanabe,H. and Sakaki,Y.
TITLE       Direct Submission
JOURNAL     Submitted (02-AUG-2001) Asao Fujiyama, The Institute of Physical
            and Chemical Research (RIKEN), Genomic Sciences Center (GSC);
            1-7-22 Suehiro-cho,Tsukumi-Ku, Yokohama, Kanagawa 230-0045, Japan
            (E-mail:chimbes@gsc.riken.go.jp, URL:http://hgp.gsc.riken.go.jp/,
            Tel:81-45-503-9111, Fax:81-45-503-9170)
            Clones are derived from the chimpanzee BAC library PTB This BAC end
            was generated during the Rad process and may have higher chance of
            clone tracking errors.
            PRIMERS
            Sequencing: M13Rev
            LIBRARY
            Vector 1 : pKS145
            R.Site 1 : SacI
            R.Site 2 : SacI
            Location/Qualifiers
            source      1. 677
                        /organism="Pan troglodytes"
                        /db_xref="taxon:9598"
                        /clone="PTB-098001.R"
                        /sex="male"
                        /cell_type="lymphoblast"
                        /clone_lib="PTB Chimpanzee Male BAC library"
BASE COUNT  178 a      175 c      176 g      147 t
ORIGIN
Query Match      16.4%; Score 283.2; DB 12; Length 677;
Best Local Similarity 48.8%; Pred. No. 2.1e-54;
Matches 294; Conservative 113; Mismatches 187; Indels 9; Gaps 3;

QY 1114 gcnngtngcnararararararararararararararararararararar 1173
      |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 68 GCTGTGCTTCAGAGAGGTAGAAGCCCAAGCCTTTGGAGCTTCATGTGCTTGAGACT 127

QY 1174 gtnngcnaararararararararararararararararararararar 1233
      |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 128 ACAGGTGCAAGAGATGACAAATTTGAGTTGTGACCTCAGCTAGATTT-AGACGATG 186

QY 1234 taygnaaycngtlytgagcmmngncaraartlycngtngnngnswngntgmg 1293
      |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 187 TTATGAAACACCTGTGATGCCAGATTTGCTCAGAGGTGAGAGGACCATCATYTGAGA 246

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